



CHEMISTRY

BOOKS - PATHFINDER CHEMISTRY (BENGALI ENGLISH)

THERMODYNAMICS

Question Bank

1. Which of the following statement is incorrect?

- A. thermodynamics deals with the relationship between heat and work
- B. thermodynamics deals with the energy changes associated with all physical and chemical processes.
- C. thermodynamics helps us to predict the direction in which a process or a chemical reaction would proceed and to what extent?
- D. thermodynamics helps us to calculate the rate of reaction

Answer: D

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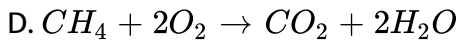
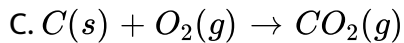
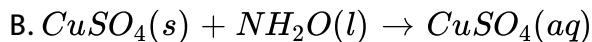
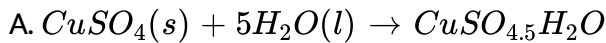
2. A system is said to be an isolated system when it is

- A. incapable of exchanging energy with the surroundings
- B. incapable of exchanging matter with the surroundings
- C. incapable of exchanging both matter and energy with the surroundings
- D. none of these

Answer: C

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3. Some reactions are given which of the following reaction refers enthalpy of hydration.



Answer: A

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4. For reaction $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ which of the following is valid

A. $\Delta H = \Delta U$

B. $\Delta h > \Delta U$

C. $\Delta H < \Delta U$

D. none of these

Answer: C

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5. Calorific values of food and fuel are determined by

- A. Bunsen's calorimeter
- B. bomb calorimeter
- C. Beckmann's thermometer
- D. none of these

Answer: B

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6. ΔH for the combustion of a compound is

- A. positive
- B. zero
- C. negative

D. may be positive or negative

Answer: C



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7. Which of the following statement is true: A system is said to be in a state of thermodynamic equilibrium when

- A. its composition remains fixed and definite
- B. the temperature at all parts of the system is the same and also identical with that of surroundings
- C. there is no unbalanced force between different parts of the system or between the system and the surroundings
- D. all are correct

Answer: D



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8. What happens when sodium cyanide solution is added to silver nitrate solution in excess?

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9. The water from the slopes of himalayas flows down the ganges into the indian ocean such natural flow of a liquid downward is

- A. non spontaneous and reversible
- B. spontaneous and reversible
- C. spontaneous and irreversible
- D. non spontaneous and irreversible

Answer: C

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10. For the combustion of heptane, Δn will be

A. (-4)

B. (-3)

C. (-2)

D. (-1)

Answer: A



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11. At 298K in a constant volume calorimeter 0.01 mole of X was detonated when 8180 calories of heat was released ΔU per mole of 'X' exploded is

A. (-808)cal/mol

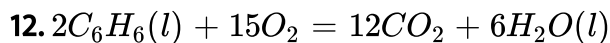
B. 880 kcal/mole

C. (-818)kcal/mole

D. (-8180)kcal/mole

Answer: C

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$2C_6H_6(g) + 15O_2 = 12CO_2 + 6H_2O(l)$ for the two reactions ΔH will be

- A. different
- B. same
- C. almost equal
- D. none of these

Answer: A

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13. Which of the following is correct one:

- A. isothermal reversible work $$'lt'$ isothermal irreversible work$
- B. isothermal reversible work = isothermal irreversible work
- C. isothermal reversible work $$'gt'$ isothermal irreversible work$
- D. none of these

Answer: C

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14. Which of the following is incorrect one:

A. Work done by the system

IUPAC (-ve)

Non IUPAC (+ve)

B. Work done on the system

IUPAC (+ve)

Non-IUPAC (-ve)

C. Heat absorbed by the system

IUPAC (+ve)

Non IUPAC (-ve)

D. Heat lost from the system

IUPAC (+ve)

Non IUPAC (-ve)

Answer: C



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15. In a reversible adiabatic change the entropy change is

A. (+ve)

B. (-ve)

C. zero

D. may be (+ve) or(-ve)

Answer: C

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16. If $H^+ + OH^- \rightarrow H_2O + 13.7kcal$ then heat of complete neutralisation of one gram mole of H_2SO_4 with a base will be

A. 13.7 k cal

B. 27.4 k cal

C. 6.85 k cal

D. 3.425 k cal

Answer: C

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17. Which of the following is an intensive property?

A. volume

B. mass

C. density

D. energy

Answer: B

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18. The unit of entropy is

A. $JK^{-1}mol^{-1}$

B. Jg^{-1}

C. $Jmol^{-1}$

D. $K^{-1}mol$

Answer: C

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19. Which of the following has highest entropy?

- A. water
- B. graphite
- C. mercury
- D. hydrogen

Answer: A

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20. For the reaction between CO_2 and graphite
 $CO_2(g) + C(s) \rightarrow 2CO(g)$, $\Delta H = +170KJ$ and $\Delta S = 170JK^{-1}$ the
reaction is spontaneous at

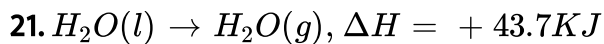
- A. 298K
- B. 500K

C. 900K

D. 1200K

Answer: D

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$H_2O(s) \rightarrow H_2O(l), \Delta h = + 6.05KJ$ the value of ΔH_{vap} or *isatation* of ice is

A. $49.75kJmol^{-1}$

B. $43.7KJmol^{-1}$

C. $37.56KJmol^{-1}$

D. none of these

Answer: D

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22. The enthalpy of formation of ammonia is $x \text{ KJmol}^{-1}$. The enthalpy of the reaction $2\text{N}_2(g) + 6\text{H}_2(g) \rightarrow 4\text{NH}_3(g)$

- A. $(-x)\text{KJ}$
- B. $x \text{ KJ}$
- C. $4x \text{ KJ}$
- D. $(-4x)\text{KJ}$

Answer: A



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23. In which of the following neutralization the heat of neutralization will be highest?

- A. NH_4OH and H_2SO_4
- B. HCl and NaOH

C. CH_3COOH and NH_4OH

D. $H_2C_2O_4$ and KOH

Answer: D

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24. Which of the following units represents largest amount of energy?

A. electron volt

B. caloric

C. joule

D. erg

Answer: B

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25. The value of ΔH for the process $NH_4OH(aq) \rightarrow NH_4^+ + Cl^-$ is

- A. negative
- B. positive
- C. zero
- D. unpredictable

Answer: B



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26. The heats of combustion of methane, carbon and hydrogen are a, b and c calories respectively what will be heat of formation of CH_4 ?

- A. $(a+c)-(a+b)$
- B. $(b+c)+(c-a)$
- C. $a+b+c$
- D. $((a+b)/2)-c$

Answer: B

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27. If an endothermic reaction is non spontaneous at freezing point of water and becomes feasible at its boiling point then

A. $\Delta H = (-ve)$, $\Delta S = (+ve)$

B. $\Delta H = (+ve)$, $\Delta S = (+ve)$

C. $\Delta H = (-ve)$, $\Delta S = (-ve)$

D. $\Delta H = (+ve)$, $\Delta S = (-ve)$

Answer: B

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28. To determine the enthalpy of neutralisation of HCl with NaOH we use

A. $0.3N HNO_3, 0.3(N) NaOH$

B. $0.4(N) HNO_3, 0.4(N) NaOH$

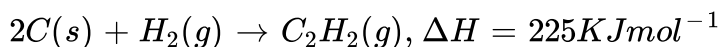
C. $1(N) HNO_3, 1(N) NaOH$

D. any of these

Answer: B

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29. Using the data provided calculate the multiple bond energy ($KJmol^{-1}$) of a C=C bond in C_2H_2 consider the bond energy of a C-H bond is $350KJmol^{-1}$



A. 1165

B. 837

C. 865

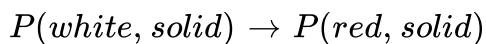
D. 815

Answer: D



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30. Which of the following for the given transformation is incorrect?



A. Allotropic change

B. ΔH is positive

C. ΔS is negative

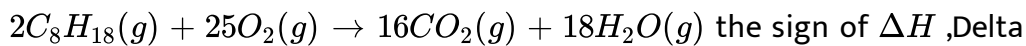
D. ΔS is positive

Answer: D



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31. For the following reaction occurring in an automobile



the sign of ΔH , ΔS and ΔG would be

A. (-),(+),(+)

B. (+),(+),(-)

C. (+),(-),(+)

D. (-),(+),(-)

Answer: D



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32. The energy for molecular motion appears in the form of

A. friction

B. potential energy

C. heat

D. temperature

Answer: D



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33. Four grams of helium gas is expanded from 1 atm to one tenth of its original pressure at 30° change in entropy assuming ideal gas behaviour is:

A. $38.3JK^{-1}$

B. $76.6JK^{-1}$

C. $19.15JK^{-1}$

D. $100JK^{-1}$

Answer: C



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34. The enthalpy of neutralisation of HCl by NaOH is -57.1 KJ and that of HCN by NaOH is $-12.1 \text{ KJ mol}^{-1}$ the enthalpy of ionization of HCN is

A. $(-69.2) \text{ KJ}$

B. $(-45) \text{ KJ}$

C. $(-69.2) \text{ KJ}$

D. 45 KJ

Answer: C



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35. Whenever an acid is neutralised by a base the net reaction is $H^+(aq) + OH^-(aq) \rightarrow H_2O(l) + 57.1 \text{ KJ}$ calculate the heat evolved in the following experiment:

0.25 mol of HCl solution is neutralised by 0.25 mol of NaOH solution amount of heat liberated for the reaction is

A. $(+57.1) \text{ KJ}$

B. (-57.1)KJ

C. (+14.3)KJ

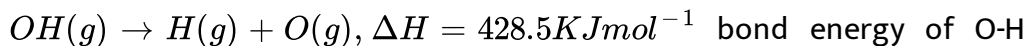
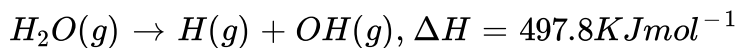
D. (-14.3)KJ

Answer: D



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36. The bond dissociation energies of two O-H bond in water are:



bond is

A. $463.15KJmol^{-1}$

B. $926.30KJmol^{-1}$

C. $231.05KJmol^{-1}$

D. $1390.45KJmol^{-1}$

Answer: C



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37. System performed pressure volume work when in between system and surrounding there is

- A. difference in temperature
- B. difference in volume
- C. difference in pressure
- D. difference in internal energy

Answer: A



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38. Change in internal energy of a system is

- A. heat absorbed by the system+work is done by the system
- B. heat liberated by the system+work is done by the system
- C. heat absorbed by the system +work is done on the system
- D. heat liberated by the system+work is done on the system

Answer: C

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39. Which of the following statement is correct?

- A. q -is a state function
- B. w -is a state function
- C. q and w are not state function together
- D. q and w are state function together

Answer: C

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40. For endothermic reaction which is correct?

A. $H_p > H_r$

B. $H_p < H_r$

C. $H_p = H_r$

D. sign of ΔH is negative

Answer: D



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41. Which of the following element standard enthalpy is not to be equal to zero?

A. C(diamond)

B. C(graphite)

C. liquid mercury

D. rhombic sulphur

Answer: B

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42. When the bond of $CH_4(g)$ is dissociated then

- A. the bond energy of all four C-H bonds is same
- B. the bond energy of two C-H bonds is same
- C. the bond energy of all four C-H bonds are different from each other
- D. the bond energy of three C-H bonds is same

Answer: A

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43. When a liquid substance is converted to solid substance its entropy

- A. increases
- B. decreases
- C. remains same
- D. any one of the above

Answer: C

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44. When ozone is converted to oxygen then entropy

- A. decreases
- B. increases
- C. equal to zero
- D. remains same

Answer: B

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45. A reaction occurs spontaneously if

- A. $\Delta H > T\Delta S$ and both ΔH and ΔS are positive
- B. $\Delta H < T\Delta S$ and ΔH is (+ve) and ΔS is (-ve)
- C. $\Delta H < T\Delta S$ and both ΔH and ΔS are positive
- D. $\Delta H = T\Delta S$ and both ΔH and ΔS are positive

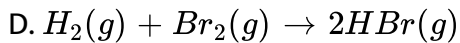
Answer: B



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46. Assume the reaction is carried out in an open system for which reaction will $\Delta H = \Delta U$?

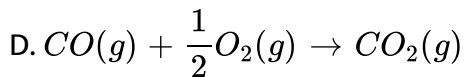
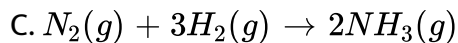
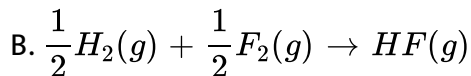
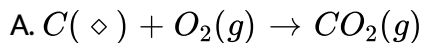
- A. $C(S) + 2H_2O(g) \rightarrow 2H_2(g) + CO_2(g)$
- B. $PCl_5(g) \rightarrow PCl_3(g) + Cl_2(g)$
- C. $2CO(g) + O_2(g) \rightarrow 2CO_2(g)$



Answer: C

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47. Which of the following reaction does not define standard enthalpy of formation (ΔH_f°)?



Answer: D

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48. Which of the following is correct ?

- A. evaporation of water causes an increase in disorder of the system
- B. melting of ice causes a decrease in randomness of the system
- C. condensation of steam causes an increase of disorder of the system
- D. there is practically no change in the randomness of the system when water is evaporated

Answer: A

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49. Write true or false of the following statements:

Molar mass is an extensive property of a system

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50. Write true or false of the following statements:

Degree of randomness increases during ice formation



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51. Write true or false of the following statements:

The internal energy of 1 mole carbon dioxide will be different from the internal energy of 1 mole sulphur dioxide even under similar conditions of temperature and pressure.



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52. Write true or false of the following statements:

No work is done during free expansion of an ideal gas under reversible or irreversible condition.



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53. Write true or false of the following statements:

A beaker containing boiling water is an example of open system.



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54. Fill in the blanks:

The two important modes of transference of energy are _____ and _____.



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55. Fill in the blanks:

For an ideal gas undergoing an _____ change the internal energy of the system does not change.



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56. Fill in the blanks:

At equilibrium the change of entropy is _____.



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57. Fill in the blanks:

At the melting point of ice ΔG is_____.



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58. Fill in the blanks:

The decrease in _____ is equal to the useful work done by the system.



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59. For a reaction both ΔH and ΔS are positive, under what condition will the reaction occur spontaneously?



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60. Write chemical equation for the enthalpy of combustion of ethanol.

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61. Name two thermodynamic properties which are additive.

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62. Ammonia is prepared as $3H_2(g) + N_2(g) \rightarrow 2NH_3(g)$ what should be the sign of ΔS for this reaction?

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63. What are the units of entropy and free energy?

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64. How is free energy change related to equilibrium constant for the reaction?

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65. If standard free energy change for a reaction is found to be zero what is its equilibrium constant?

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66. $Cl_2(g) \rightarrow 2Cl(g) \Delta H^\circ = 242 KJmol^{-1}$ what is its enthalpy of atomization?

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67. Write down the statement of second law of thermodynamics.

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68. which of the following has greater fuel efficiency?

Hydrogen, methane, LPG

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69. Write the mathematical formulation of the first law of thermodynamics.

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70. Give an example of a process where $q=0$

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71. under what condition heat of reaction of a chemical reaction becomes equal to internal energy change?

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72. Among the enthalpy of solution and enthalpy of hydration .which is always an exothermic process?



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73. Give an example of a spontaneous process where the randomness of the system decreases.



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74. Which has the more entropy?

1 mole H_2 gas ($T=300K$, $V=5mL$)

1 mole H_2 gas ($T=300K$, $V=10mL$)



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75. Which has the more entropy?

1 mole H_2 gas ($T=300K, V=10L$)

1 mole H_2 gas ($T=300K, V=30L$)



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76. When $\Delta G = 0$ the process will be at _____



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77. At what temperature entropy of a perfect crystalline substance is taken as zero?



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78. For the reaction $x + y \rightarrow p + q$ the value of ΔH is +a K cal what is the value of ΔH for the reaction $p + q \rightarrow x + y$?

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79. What happens when Mercuric iodide reacts with potassium iodide?

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80. When both ΔH and ΔS become (-ve) the process will be spontaneous at ___ tempratures.

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81. Which factor does remain constant in an isobaric process?

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82. Write the expression of Gibbs Helmholtz equation

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83. When is $\Delta S = \frac{\Delta}{T}$

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84. State first law of thermodynamics.

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85. A system absorbs 115J heat from the surroundings and 290 J work is done on the system calculate the change in internal energy of the system.

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86. Define system and surrounding.

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87. A gas absorbs 120J of heat and expands against the external pressure of 1.10 atm from a volume of 0.5 L to 2L what is the change in internal energy?(1 L atm=101.3J)

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88. What is enthalpy of a system?

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89. Show that at constant volume $\Delta U = q_v$

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90. Write one limitation of the first law of thermodynamics .

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91. The latent heat of evaporation of Bromine at $59^{\circ}C$ at 1 atm is 29.2 KJ/mol what are ΔH and ΔU of this process?

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92. Explain the term state function of a system

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93. for the reaction $2Cl(g) \rightarrow Cl_2(g)$ What are the signs of ΔH and ΔS ?

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94. To increase the temperature of 31.75 mm of Cu from $23^{\circ}C$ to $28^{\circ}C$,50 calorie of heat is required find heat capacity of copper

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95. To increase the temperature of 31.75 mm of Cu from $23^{\circ}C$ to $28^{\circ}C$, 50 calorie of heat is required find heat capacity of copper

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96. To increase the temperature of 31.75 mm of Cu from $23^{\circ}C$ to $28^{\circ}C$, 50 calorie of heat is required find heat capacity of copper

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97. Which of the following are not state function?
Entropy, temperature, heat, enthalpy, internal energy, work

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98. At 25°C What is the difference between ΔH and ΔS for the following reaction? $2\text{H}_2(\text{g}) + \text{O}_2(\text{g}) \rightarrow 2\text{H}_2\text{O}(\text{l})$

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99. Establish the relation between ΔH and ΔU for a gaseous system at constant temperature and pressure

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100. $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightarrow 2\text{NH}_3(\text{g}) \Delta H_r^{\circ} = -92.4\text{KJ}$

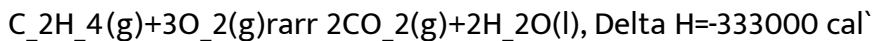
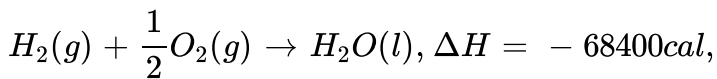
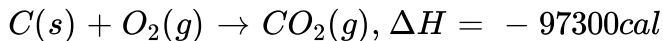
What is the standard enthalpy of formation of NH_3 ?

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101. Write Hess's law.

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102. Using the data given calculate the heat of formation of ethylene.

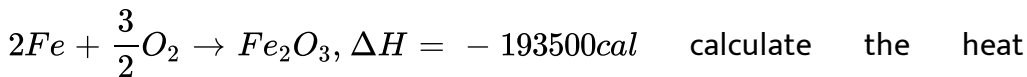
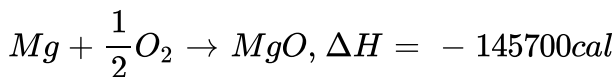


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103. Write the thermodynamic relation generally used to predict a reaction is spontaneous or not

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104. The heats of oxidation of Mg and Fe are given:



produced in the reaction $3Mg + Fe_2O_3 = 3MgO + 2Fe$

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105. For the reaction at 298 K, $2A + B \rightleftharpoons C$, $\Delta H = 400 \text{ KJ mol}^{-1}$ and $\Delta S = 0.2 \text{ KJ K}^{-1} \text{ mol}^{-1}$. At what temperature will the reaction become spontaneous?

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106. The equilibrium constant for a reaction is 10 what will be the value of ΔG° ? [$R = 8.314 \text{ JK}^{-1} \text{ mol}^{-1}$, $T = 300 \text{ K}$]

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107. Diamond is an elementary substance yet its standard heat of formation is not taken as zero why?

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108. The enthalpy change for the transition of liquid water to steam is 40.8 KJ mol^{-1} at 373 K calculate the entropy of evaporation of water.

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109. 10 gm argon is compressed isothermally and reversibly at 27°C from 10 L to 5L calculate q, w and ΔU for the process [$R=2 \text{ cal K}^{-1} \text{ mol}^{-1}$] At wt.of Ar=40]

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110. Enthalpy of neutralisation of any strong acid with a strong base is always constant.why so?

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111. Write down the dimension of entropy.





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112. Write the mathematical form of first law of thermodynamics for the following processes Adiabatic



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113. Write the mathematical form of first law of thermodynamics for the following processes isothermal



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114. Latent heat of fusion of tin is 14 cal/g calculate the change in entropy when 10 gm of tin is melted at 232°C ,



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115. Enthalpies of solution of $(BaCl_2) \cdot (2H_2O)$ and $BaCl_2$ are 8.8 and $(-20.6) KJmol^{-1}$ respectively calculate the heat of hydration of for conversion of $BaCl_2$ to $(BaCl_2) \cdot (2H_2O)$

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116. Which of the following state of bromine will have standard heat of formation to be equal to zero liquid bromine, solid bromine, gaseous bromine molecules.

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117. Calculate ΔH_r° for the reaction $C_2H_4 + 3O_2 \rightarrow 2CO_2 + 2H_2O$ the average bond enthalpies of various bonds are

Bond $C - H$ $O = O$ $C = O$ $H - O - H$ $C - C$ Bond enthalpy $KJ \text{ mol}^{-1}$

414, 499, 724, 460, 619

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118. Heat of neutralisation of reaction between weak acid and weak base is less than that of reaction between strong base.explain why?

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119. Calculate the enthalpy of sublimation of Na metal from the following data given $\Delta H_{fus}^{\circ} = 2.6KJmol^{-1}$

$$\Delta H_{vap}^{\circ} = 98KJmol^{-1}$$

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120. Enthalpies of formation of $CO(g)$, $CO_2(g)$, $N_2O(g)$ and $N_2O_4(g)$ are -110,-393,81 and $9.7KJmol^{-1}$ respectively calculate ΔH_r for the reaction
$$:N_2O_4(g) + 3CO(g) \rightarrow N_2O(g) + 3CO_2(g)$$

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121. The ΔH_r° for the reaction $4S(s) + 6O_2(g) \rightarrow 4SO_3(g)$ is -1583.2 KJ
calculate ΔH_f° of sulphur trioxide.

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122. Standard entropy of X_2 , Y_2 and XY_3 are $60, 40$, and $50 \text{ J K}^{-1} \text{ mol}^{-1}$
respectively for the reaction $\frac{1}{2}X_2 + \frac{3}{2}Y_2 \rightarrow XY_3$, $\Delta H = -30 \text{ KJ}$
what will be the temperature at equilibrium?

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123. What happens when phenol reacts with zinc dust?

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124. For a spontaneous reaction $\Delta H = (-ve)$, $k > 1$ and $E_{cell}^\circ = +ve$
are these informations correct or not?



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125. The bond dissociation energies of XY , X_2 and Y_2 (all diatomic molecules) are in the reaction 1:1:0:5 and ΔH_r for the formation of XY is -200 KJmol^{-1} what will be the bond dissociation energy of X_2 ?



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126. Calculate the free energy change and entropy change per mole when liquid water boils at 1 atmosphere latent heat of vaporization of water = $2.0723 \text{ KJmol}^{-1}$



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127. What is the entropy change for dissolving 1 mole of NaCl at 298 K
lattice energy =

$+777.8 \text{ KJmol}^{-1}$ $\Delta H_{\text{Hydration of NaCl}} = -774.1 \text{ KJmol}^{-1}$, $\Delta G_{\text{dissolution of}}$



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128. Calculate the entropy change involved in the isothermal expansion of 5 mol of ideal gas from a volume of 10 L to 100 L at 300 K

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129. What is internal energy?

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130. If the internal energy of the system is decreased by 100 joule and heat released by the system is 80 joule then calculate work done by the system.

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131. Standard heat of formation of $C_2H_6(g)$, $CO_2(g)$ and $H_2O(l)$ are -21.1 , -94.1 and $-68.3 Kcal mol^{-1}$ respectively calculate standard heat of combustion of $C_2H_6(g)$

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132. Define enthalpy of transition.

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133. Heat of combustion of rhombic and monoclinic sulphur be $-295.1 KJ mol^{-1}$ and $-297.4 KJ mol^{-1}$ respectively calculate the enthalpy of transition from monoclinic sulphur to rhombic sulphur

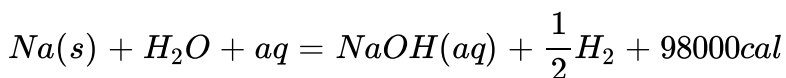
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134. Which is correct? (i) $H_{charcoal}^{\circ} = 0$ (ii) $H_{graphite}^{\circ} = 0$



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135. From the following data calculate the heat of formation of caustic soda (NaOH)



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136. Explain how Bakelite is formed from phenol?



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137. What is the monomer of *Nylon – 6*?



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138. Define Bond energy



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139. The bond energy of $C - C$, $C = C$, $H-H$ and $C-H$ bond are 350, 600, 400 and 410 KJmol^{-1} respectively calculate the change in enthalpy of the hydrogenation reaction of ethylene.

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140. Give example of the processes where the change in entropy will be equal to zero.

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141. under what condition an adiabatic process may become isentropic?

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142. Write down the statement of third law of thermodynamics.



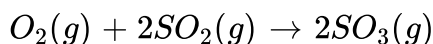
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143. Calculate the standard entropy change of the following reaction at $25^\circ C$: $SO_2(g) + \frac{1}{2}O_2(g) \rightarrow SO_3(g)$ at $25^\circ C$ tempr.and 1 atm pressure the standard entropy of $SO_2(g)$, $O_2(g)$ and $SO_3(g)$ are 248.5,205.0 and $256.2JK^{-1}mol^{-1}$ respectively



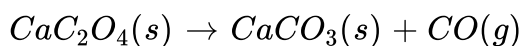
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144. Predict the sign of entropy change is the following reactions



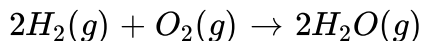
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145. Predict the sign of entropy change is the following reactions



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146. Predict the sign of entropy change is the following reactions



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147. Heat capacity (C_p) is an extensive property but specific heat (C) is an intensive property. What will be the relation between C_p and C for 1 mole of water ?

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148. Calculate Δq , W , ΔE for the isothermal reversible expansion of one mole of an ideal gas from an initial pressure of 1.0 bar to final pressure of 0.1 bar at a constant temperature of 273 K.

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149. 1 mol of an ideal gas undergoes reversible isothermal expansion from an initial volume V_1 to final volume $10 V_1$ and does 10 kJ of work. The initial pressure was 1×10^7 Pa, Calculate the value of V_1 .

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150. Calculate the amount of work done during the process when one mole of an ideal gas is allowed to expand freely into vacuum.

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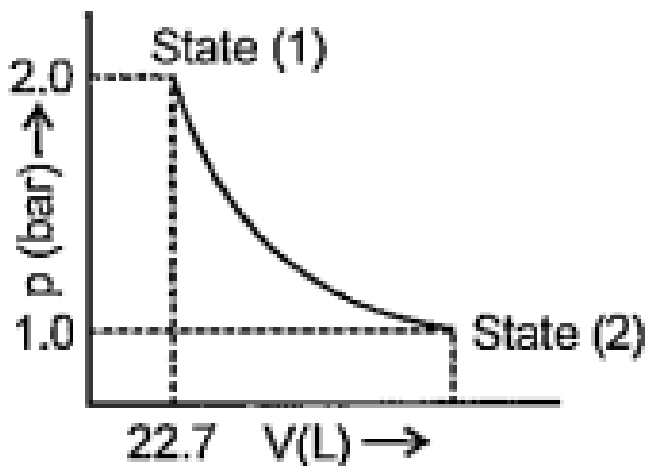
151. Two litres of an ideal gas at a pressure of 10 atm expands isothermally into a vacuum until its total volume is 10 litres. How much heat is absorbed & how much work is done in the expansion ?

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152. Two litres of an ideal gas at a pressure of 10 atm expands isothermally into a vacuum until final volume is of 10 litres conducted reversibly. How much heat is absorbed ?

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153. 1.0 mole of a monoatomic ideal gas is expanded from state (1) to state (2) as shown in figure. Calculate the work done for the expansion of gas from state (1) to state (2) at 298 K



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154. One mole of an ideal gas at STP is heated to $27^{\circ}C$ and compressed to 1.5 atm. Calculate ΔE and ΔH for the change in state. Assume $C_v = \frac{3}{2}R$ for the ideal gas.

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155. Calculate heat of the following reaction at constant pressure, $F_2O(g) + H_2O(g) \rightarrow O_2(g) + 2HF(g)$ The heats of formation (in Kcal mol^{-1}) of $F_2O(g)$, $H_2O(g)$ and $HF(g)$ are 5.5, -57.8 and -64.2 respectively.

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156. If water vapour is assumed to be a perfect gas and molar enthalpy change for vapourisation of 1 mole of water at 1 bar & $100^{\circ}C$ is $41 \text{ kJ } mol^{-1}$, calculate the internal energy change when 1 mol of water is vapourised at 1 bar pressure & $100^{\circ}C$

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157. If water vapour is assumed to be a perfect gas and molar enthalpy change for vapourisation of 1 mole of water at 1 bar & $100^{\circ}C$ is 41 kJ mol^{-1} , calculate the intenal energy change when 1 mol of water is converted into ice.

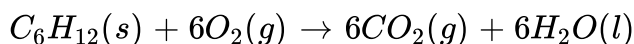
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158. The enthalpy of combustion of methane, graphite & dihydrogen at 298 K are, $-890.3 \text{ kJ mol}^{-1}$, $393.5 \text{ kJ mol}^{-1}$ & $285.8 \text{ kJ mol}^{-1}$, respectively.

Enthalpy of formation of $CH_4(g)$ will be :

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159. Heat of reaction for,



at constant pressure is -651 kcal at $17^{\circ}C$. Calculate the heat of reaction at constant volume at $17^{\circ}C$.

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160. Calculate the number of kJ of heat necessary to raise the temperature of 60.0 g of aluminium from 35°C to 55°C . Molar heat capacity of Al is $24\text{Jmol}^{-1}\text{K}^{-1}$. Molar mass of Al = 27gmol^{-1} .

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161. Enthalpies of formation of $\text{CO}(g)$, $\text{CO}_2(g)$, $\text{N}_2\text{O}(g)$ and $\text{N}_2\text{O}_4(g)$ are -110, -393, 81 and 9.7kJmol^{-1} respectively calculate ΔH_r for the reaction

$$:\text{N}_2\text{O}_4(g) + 3\text{CO}(g) \rightarrow \text{N}_2\text{O}(g) + 3\text{CO}_2(g)$$

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162. Calculate the enthalpy change on freezing of 1.0 mole of water at 10°C to ice at -10°C , $\Delta_{fus} H = 6.03\text{kJmol}^{-1}$ at 0°C ,

$$C_p[\text{H}_2\text{O}(l)] = 75.3\text{Jmol}^{-1}\text{K}^{-1}$$

$$C_p[\text{H}_2\text{O}(s)] = 36.8\text{Jmol}^{-1}\text{K}^{-1}$$



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163. Calculate the enthalpy change for the process :

$CCl_4(g) \rightarrow C(g) + 4Cl(g)$ and calculate the bond enthalpy of $C - Cl$ in

$CCl_4(g)$

$$\Delta (vap) H^\circ (CCl_4) = 30.5 \text{ kJ mol}^{-1}$$

$$\Delta_f H^\circ (CCl_4) = -135.5 \text{ kJ mol}^{-1}$$

$$\Delta_a H^\circ (C) = 715.0 \text{ kJ mol}^{-1}$$

$$\Delta_a H^\circ (Cl_2) (\text{atomisation}) = 242 \text{ kJ mol}^{-1}$$

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164. For an isolated system, $\Delta U = 0$, what will be ΔS ?

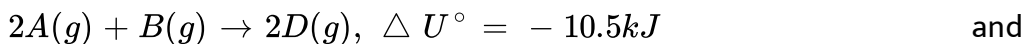
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165. Calculate the entropy change in surroundings when 1.00 mole of $H_2O(l)$ is formed under standard conditions.

$$\Delta_f H^\circ = 286 \text{ kJ mol}^{-1}.$$

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166. For the reaction



$$\Delta S^\circ = -44.1 \text{ JK}^{-1}$$

Calculate ΔG° for the reaction, and predict whether the reaction may occur spontaneously.

$$(R = 8.314 \times 10^{-3} \text{ kJK}^{-1} \text{ mol}^{-1}, T = 298 \text{ K})$$

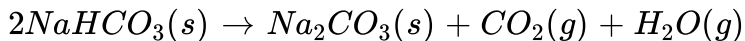
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167. Predict in the given question, entropy increases/ decreases

A liquid crystalizes into a solid

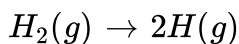
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168. Predict in the given question, entropy increases/ decreases



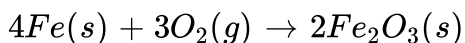
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169. Predict in the given question, entropy increases/ decreases



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170. For oxidation of Fe



entropy change is $-549.4\text{JK}^{-1}\text{mol}^{-1}$ at 298K. In spite of negative entropy change of this reaction, why is the reaction spontaneous ?

$$\Delta H^\circ = -1648 \times 10^3 \text{Jmol}^{-1}$$

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171. For the reaction at 298K



$$\Delta H = 400 \text{kJmol}^{-1}, \Delta S = 0.2 \text{kJK}^{-1}\text{mol}^{-1}$$

At what temperature will the reaction become spontaneous considering

ΔH & ΔS to be constant over the temperature range.



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172. If the internal energy of an ideal gas decreases by the same amount as the work done by the system, the process is

- A. cyclic
- B. isothermal
- C. adiabatic
- D. isolated

Answer: C



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173. In which of the following pairs, both properties are intensive ?

A. Pressure, temperature

B. Density, volume

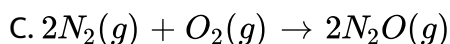
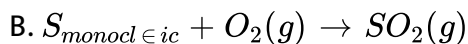
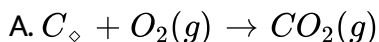
C. Temperature, density

D. Pressure, volume

Answer: C

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174. Which among the following represents the reaction of formation of the product ?

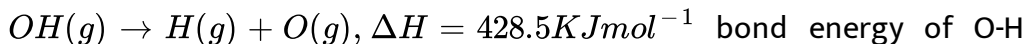
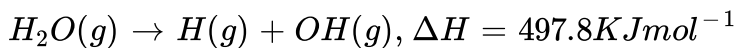


D. None of the above

Answer: D

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175. The bond dissociation energies of two O-H bond in water are:



bond is

A. 498 kJ mol^{-1}

B. 463 kJ mol^{-1}

C. 428 kJ mol^{-1}

D. 70 kJ mol^{-1}

Answer: B

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176. For an ideal gas, the Joule Thomson coefficient is

- A. zero
- B. positive
- C. negative
- D. depends on atomicity of gas

Answer: A



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177. If 150 kJ of energy is needed for muscular work to walk a distance of 1 km, then how much of glucose one has to consume to walk a distance of 5 km, provided only 30% of energy is available for muscular work. The enthalpy of combustion of glucose is 3000kJmol^{-1}

- A. 75 g
- B. 30 g

C. 180 g

D. 150 g

Answer: D

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178. For a particular reaction, $\Delta H^\circ = -38.3 \text{ kJ mol}^{-1}$ and $\Delta S^\circ = -113 \text{ JK}^{-1} \text{ mol}^{-1}$. This reaction is

A. spontaneous at all temperatures

B. non-spontaneous at all temperatures

C. spontaneous at temperature below 66° C

D. spontaneotis at temperature above 66° C

Answer: C

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179. For a process to be in equilibrium, it is necessary that

A. $\Delta S_{system} = \Delta S_{surr}$

B. $\Delta S_{system} = - \Delta S_{surr}$

C. $\Delta S_{system} = 0$

D. $\Delta S_{surr} = 0$

Answer: B



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180. Which of the following statement(s) must be true for the entropy of a pure solid to be zero ?

I. The temperature must be zero kelvin

II. The solid must be perfectly crystalline

III. The solid must be an element

IV. The solid must be ionic

A. I

B. I AND II

C. I, II AND III

D. ALL ARE CORRECT

Answer: B

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181. Which of the following conditions may lead to a non-spontaneous change ?

A. $\Delta H = +ve$, $\Delta S = -ve$

B. $\Delta H = -ve$, $\Delta S = +ve$

C. $\Delta H = -ve$, $\Delta S = -ve$

D. $\Delta H = +ve$, $\Delta S = +ve$

Answer: A

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182. For the process, $NH_3(g) + HCl(g) \rightarrow NH_4Cl(s)$

- A. $\Delta H = +ve, \Delta S = ve+$
- B. $\Delta H = -ve, \Delta S = +ve$
- C. $\Delta H = +ve, \Delta S = -ve$
- D. $\Delta H = -ve, \Delta S = -ve$

Answer: D



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183. Which of the following statements is false ?

- A. Work is a state function
- B. Temperature is a state function
- C. Change in the state is completely defined when the initial and final states are specified

D. Work appears at the boundary of the system

Answer: A



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184. In thermodynamics, a process is called reversible when

- A. the surroundings and system change into each other
- B. there is no boundary between system and surroundings
- C. the surroundings are always in equilibrium with the system
- D. the system charges into the surroundings spontaneously

Answer: C



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185. Which of the following is not correct for adiabatic process ?

A. $TV^{\gamma-1} = k$

B. $PV^{\gamma} = K$

C. $\gamma = \frac{C_v}{C_p}$

D. $PT^{\frac{\gamma}{1-\gamma}} = K$

Answer: C

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186. A piston filled with 0.04mole of an ideal gas expands reversibly from 50.0mL to 375mL at a constant temperature of $37.0^{\circ}C$. As it does so, it absorbs 208J of heat. The values of q and W for the process will be ($R = 8.314\text{J/mol K}$, $\ln 7.5 = 2.01$)

A. $q = -208J, w = +208J$

B. $q = +208J, w = +208J$

C. $q = +208J, w = -208J$

D. $q = -208J, w = -208J$

Answer: C



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187. Temperature of an ideal gas increases in

- A. adiabatic expansion
- B. adiabatic compression
- C. isothermal expansion
- D. isobaric expansion

Answer: B



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188. Which of the following statements is incorrect

- A. The specific heat capacity of substance is greater in the solid state than in liquid state.
- B. The specific heat capacity of a substance is greater in gaseous state than in the liquid state
- C. The latent heat of vaporization of a substance is greater than that of fusion.
- D. The internal energy of an ideal gas is a function of its temperature.

Answer: B

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189. Which is equal to total work done ?

A. $P \Delta V$

B. ΔG

C. ΔH

D. ΔE

Answer: A



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190. $\left(\frac{\delta\Delta H}{\delta P}\right)_T$ for an ideal gas is equal to

A. $\Delta V \frac{RT}{P}$

B. $T\Delta V \frac{R}{P}$

C. zero

D. $\left[\left(\delta \frac{\Delta V}{\delta T}\right)\right]$

Answer: C



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191. The standard enthalpy of formation of $H_2O(g)$ at 298 K is $-241.82 \text{ kJ mol}^{-1}$, Calculate ΔH° at 373 K. Assume that C_p is independent of temperature

$$C_p^\circ \text{ of } H_2O(g) = 33.58 \text{ JK}^{-1} \text{ mol}^{-1}$$

$$C_p^\circ \text{ of } H_2(g) = 28.84 \text{ JK}^{-1} \text{ mol}^{-1}$$

$$C_p^\circ \text{ of } O_2(g) = 29.37 \text{ JK}^{-1} \text{ mol}^{-1}$$

- A. -242 kJ
- B. +242.6 kJ
- C. +24.26 kJ
- D. -242.6 J

Answer: A



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192. Which of the following compounds has a positive enthalpy of solution?

A. LiF

B. LiCl

C. LiBr

D. LiI

Answer: D



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193. When a bomb calorimeter is used to determine the heat of reaction, which property of the system under investigation is most likely to remain constant ?

A. Number of molecules

B. Pressure

C. Temperature

D. Volume

Answer: D



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194. The heat of combustion of benzoic acid at constant volume is -321.3 kJ at 27°C . The heat of combustion at constant pressure is

A. -321.3 kJ

B. -317.54 kJ

C. $+322.54$ kJ

D. $+321.3$ kJ

Answer: B



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195. The molar heats of combustion of $\text{C}_2\text{H}_2(\text{g})$, $\text{C}(\text{graphite})$ and $\text{H}_2(\text{g})$ are -310.62 , -94.05 , -68.32 kcal respectively. The heat of formation of

acetylene is

- A. 54.2 kcal
- B. -54.2 kcal
- C. -68.32 kcal
- D. +68.32 kcal

Answer: B



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196. The standard molar heat of formation of ethane, CO_2 and $H_2O(l)$ are respectively -21.1, -94.1 and -68.3 kcal. The standard molar heat of combustion (in kcal) of ethane will be

- A. -372
- B. 162
- C. -240

D. 183.5

Answer: A

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197. The standard enthalpies of formation of $CO_2(g)$, $H_2O(l)$ and glucose (s) at $25^\circ C$ are -400 kJ/mol, -300 kJ/mol and -1300 kJ/mol, respectively. The standard enthalpy of combustion per gram of glucose at $25^\circ C$ is

A. $+2900kJ$

B. -2900 kJ

C. -16.11 kJ

D. $+16.11kJ$

Answer: C

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198. When 1g of anhydrous oxalic acid is burnt at $25^{\circ}C$, the amount of heat liberated is 2.835 kJ. ΔH_{comb} is

- A. 255.15 kJ
- B. 445.65 kJ
- C. -255.15 kJ
- D. 155.16 kJ

Answer: C



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199. Calculate the enthalpy change when 50 mL of 0.01 M $Ca(OH)_2$ reacts with 25 mL of 0.01 M HCl. Given that ΔH_N° , of a strong acid and strong base is 57.1 kJ.

- A. 14.275 kJ
- B. 14.275 J

C. $14.275 \times 10^3 \text{ kJ}$

D. $14.375 \times 10^3 \text{ J}$

Answer: B

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200. The enthalpy of neutralization of a strong acid by a strong base is $-57.32 \text{ kJ eqv}^{-1}$. The enthalpy of formation of water is $-285.84 \text{ kJ eqv}^{-1}$.

The enthalpy of formation of hydroxyl ion is

A. $+228.52 \text{ kJ mol}^{-1}$

B. $-114.26 \text{ kJ mol}^{-1}$

C. $-288.52 \text{ kJ mol}^{-1}$

D. $+114.2 \text{ kJ mol}^{-1}$

Answer: C

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201. The enthalpy of solution of $BaCl_2(s)$ and $BaCl_2 \cdot 2H_2O(s)$ are $-20.6 kJ mol^{-1}$ and $8.8 kJ mol^{-1}$. The enthalpy change for the formation of hydrated salt is

- A. 29.8 kJ
- B. -11.8 kJ
- C. -20.6 kJ
- D. -29.4 kJ

Answer: D

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202. The heat of solution of anhydrous $CuSO_4$ is -15.9 kcal and that of $CuSO_4 \cdot 5H_2O$ is 2.8 kcal. the heat of hydration of $CuSO_4$ is

- A. $+18.7 kcal$

B. -18.7 kcal

C. -15.9 kcal

D. -2.8 kcal

Answer: B

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203. The heat of atomisation of $PH_3(g)$ is 228kcalmol^{-1} and that of $P_2H_4(g)$ is 355kcalmol^{-1} . The energy of P-P bond is

A. 62 kcal

B. 51 kcal

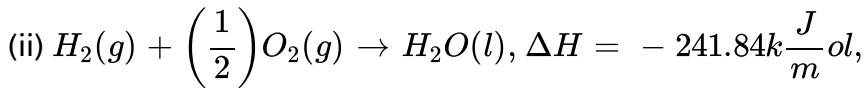
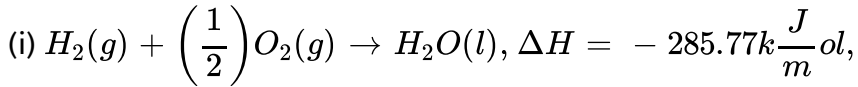
C. 52 kcal

D. 53 kcal

Answer: B

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204. The enthalpy of vapourisation of liquid water using the data:



A. +43.93

B. -43.93

C. +527.61

D. -527.61

Answer: A



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205. In reversible adiabatic process ΔS is

A. infinity

B. zero

C. equal to $C_v dT$

D. equal to $nR \frac{\ln V_2}{V_1}$

Answer: B

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206. For which of these processes is the value of ΔS negative

I. Sugar is dissolved in water.

II. Steam condenses on a surface

III. $CaCO_3$ is decomposed into CaO and CO_2

A. I ONLY

B. II ONLY

C. I AND III ONLY

D. II AND III ONLY

Answer: B

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207. The entropy change accompanying the heating of one mole of helium gas, assuming ideal behaviour from a temperature of 300 K to a temperature of 1000 K at constant pressure.

A. $25.17JK^{-1}mol^{-1}$

B. $20kJK^{-1}$

C. $2.517JK^{-1}mol^{-1}$

D. $0.2517JK^{-1}mol^{-1}$

Answer: A

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208. A reaction for which $\Delta H = -11.7 \times 10^3 Jmol^{-1}$ and $\Delta S = 105JK^{-1}mol^{-1}$ would be spontaneous when temperature is

A. equal to $111.4^\circ C$

B. equal to 111.4 K

C. $> 111.4K$

D. $< 111.4K$

Answer: D

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209. A gas expands adiabatically at constant pressure such that:

$$T \propto \frac{1}{\sqrt{V}}$$

The value of γ , i.e., $\left(\frac{C_p}{C_v}\right)$ of the gas will be

A. 1.3

B. 1.5

C. 1.7

D. 2

Answer: B

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210. One mole of an ideal gas expands reversibly and adiabatically from a temperature of 27°C . If the work done during the process is 3 kJ, then final temperature of the gas is ($C_V = 20\frac{J}{K}$)

A. 100K

B. 150K

C. 195K

D. 255K

Answer: B

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211. Enthalpy of formation of HF and HCl are -161 kJ mol^{-1} and -92 kJ mol^{-1} respectively. Which of the following statements is incorrect?

A. HCl is more stable than HF

B. HF and HCl are exothermic compounds.

C. HF is more stable than HCl.

D. The affinity of fluorine to hydrogen is greater than the affinity of chlorine to hydrogen

Answer: A

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212. The standard enthalpy of formation of NH_3 is $46kJmol^{-1}$. If the enthalpy of formation of H_2 from its atoms is $-436kJmol^{-1}$ and that of nitrogen is $-712kJmol^{-1}$, the average bond enthalpy of N-H bond in NH_3 is

A. $+1056kJmol^{-1}$

B. $-1102kJmol^{-1}$

C. $-964kJmol^{-1}$

D. $+352\text{kJmol}^{-1}$

Answer: D

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213. The heat liberated when 1.89 g of benzoic acid is burnt in a bomb calorimeter at 25°C increase the temperature of 18.94 kg of water by 0.632°C . If the specific heat of water at 25°C is 0.998 cal/g-deg , the value of the heat of combustion of benzoic acid is

A. 881.1 cal

B. 771.4 cal

C. 981.1 cal

D. 871.2 cal

Answer: B

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214. Which of the following thermodynamic relation is correct ?

A. $dG = VdP - SdT$

B. $dU = PdV + TdS$

C. $dH = TdS - VdP$

D. $dG = VdP + SdT$

Answer: A



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215. Standard entropy of X_2 , Y_2 and XY_3 are 60, 40, and $50 \text{ J K}^{-1} \text{ mol}^{-1}$ respectively for the reaction $\frac{1}{2}X_2 + \frac{3}{2}Y_2 \rightarrow XY_3$, $\Delta H = -30 \text{ KJ}$ what will be the temperature at equilibrium?

A. 750 K

B. 1000 K

C. 1250 K

D. 500 K

Answer: A

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216. Assuming ΔH° and S° do not change with temperature, calculate the boiling point of liquid 'A' using the thermodynamic data given below :

Thermodynamic data	A (liq)	A (gas)
ΔH° (kJ/mol)	-130	-100
S° ($\text{JK}^{-1} \text{mol}^{-1}$)	100	200

A. 300 K

B. 130 K

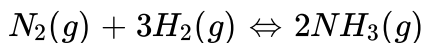
C. 150 K

D. 50 K

Answer: A

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217. The Haber's process for production of NH_3 involves the reaction :



Assuming ΔH° and ΔS° for the reaction do not change with temperature, which of the statements is true? ($\Delta H^\circ = -95\text{kJ}$ and $\Delta S^\circ = -190\text{JK}^{-1}$)

- A. Ammonia dissociates spontaneously below 500 K
- B. Ammonia dissociates spontaneously above 500 K
- C. Ammonia dissociates at all temperatures
- D. Ammonia does not dissociate at any temperature

Answer: B



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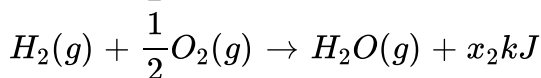
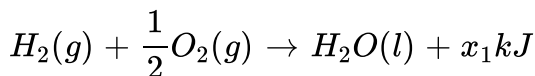
218. Which is/are the correct relationship(s) ?

- A. $\left[\frac{dE}{dT} \right]_v \neq 0$ (for ideal gas)
- B. $\left[\frac{dE}{dT} \right]_T = 0$ (for ideal gas)
- C. $\left[\frac{dV}{dT} \right]_P = \frac{nR}{P}$ (for ideal gas)
- D. $\left(\frac{\delta G}{\delta T} \right) = P$

Answer: B::C

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219. For the two equations given below :



Select the correct answer(s)

- A. $x_1 > x_2$
- B. $x_1 - x_2 > 0$
- C. $x_1 = x_2$
- D. $x_1 + x_2 = 0$

Answer: A::B



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220. $\Delta E = 0$, for which process(s)?

- A. Cyclic process
- B. Isothermal expansion
- C. Isochoric process
- D. Adiabatic process

Answer: A::B



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221. For a reaction to be spontaneous in neither direction, which of the following is/are correct regarding the closed system ?

$$(\Delta G)_{T,P} = 0$$

$$(\Delta G)_{T,P} < 0$$

$$(\Delta S)_{U,V} = 0$$

$$(\Delta S)_{U,V} > 0$$

A. A,B AND C ARE CORRECT

B. A IS CORRECT

C. B AND D ARE CORRECT

D. A and C IS CORRECT

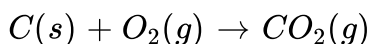
Answer: B::D



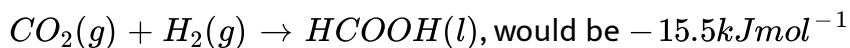
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222. The standard enthalpies of formation of $CO_2(g)$ and $HCOOH(l)$ are $-393.7kJmol^{-1}$ and $-409.2kJmol^{-1}$ respectively. Which of the following statements are correct?

A. $-393.7kJmol^{-1}$ is the enthalpy change for the reaction,



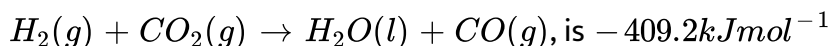
B. The enthalpy change for the reaction,



C. The enthalpy change for the reaction,



D. The enthalpy change for the reaction,



Answer: A::B



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223. Which of the following is a state function as well as an extensive property ?

A. Temperature

B. Internal energy

C. Entropy

D. Heat capacity

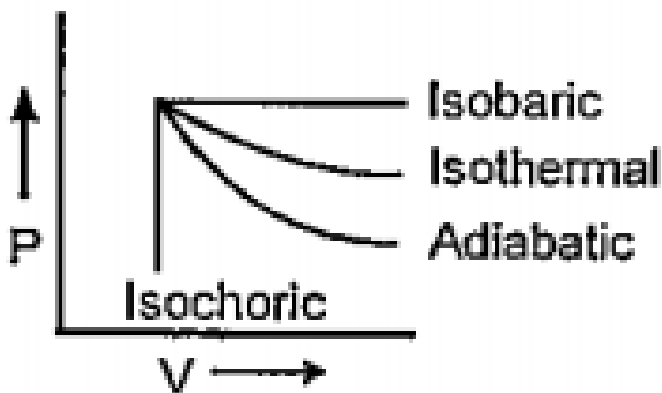
Answer: B::C::D

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224. If the system involves gaseous substance, then pressure volume work

$W_{pv} = -P_{ext}\Delta V$. It has been observed that,

$$W_{rev} > W_{ir.}$$



Also the work of isothermal & adiabatic processes are different from each other :

$$W_{isothermalreversib\le} = 2.303nRT \log\left(\frac{v_2}{v_1}\right)$$

$$W_{adiabaticreversib\le} = -C_v(T_2 - T_1)$$

If W_1, W_2, W_3, W_4 are work done in isothermal, adiabatic, isobaric & isochoric reversible processes, then correct sequence would be

- A. $W_1 > W_2 > W_3 > W_4$
- B. $W_3 > W_2 > W_1 > W_4$
- C. $W_3 > W_2 > W_4 > W_1$
- D. $W_3 > W_1 > W_2 > W_4$

Answer: D

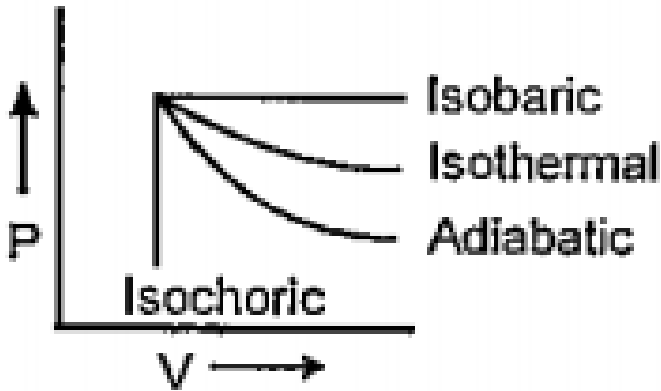


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225. If the system involves gaseous substance, then pressure volume work

$W_{pv} = -P_{ext}\Delta V$. It has been observed that,

$$W_{rev} > W_{ir}$$

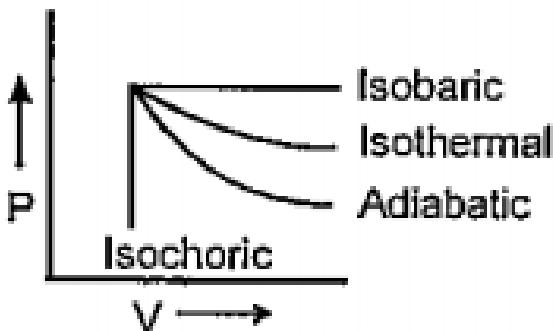


Also the work of isothermal & adiabatic processes are different from each other :

$$W_{isothermalreversib\le} = 2.303nRT \log\left(\frac{v_2}{v_1}\right)$$

$$W_{adiabaticreversib\le} = - C_v(T_2 - T_1)$$

P-V plot for two gases during adiabatic processes are given as follows :



Curves A&B should correspond to respectively

A. $He \& O_2$

B. $He \& Ar$

C. $O_2 \& He$

D. $O_2 \& F_2$

Answer: C

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226. ΔG of the system alone provides a criterion for the spontaneity of a process at constant temperature & pressure,

$\Delta G_{system} < 0$ (spontaneous)

$\Delta G_{system} = 0$ (equilibrium) $\Delta G_{system} > 0$

{(non-spontaneous) $\Delta G_{system} > 0$

is related \rightarrow the equilibrium constant as follows: $\Delta G^{\circ} = -2.303 RT$

$\log K_e$ or a spontaneous reaction ΔG and K_e will be respectively

A. -ve, +ve

B. +ve, -ve

C. zero, zero

D. -ve, -ve

Answer: A

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227. ΔG of the system alone provides a criterion for the spontaneity of a process at constant temperature & pressure,

$\Delta G_{\text{system}} < 0$ (spontaneous)

$\Delta G_{\text{system}} = 0$ (equilibrium)

$\Delta G_{\text{system}} > 0$ (non-spontaneous)

Again ΔG° is related to the equilibrium constant as follows :

$$\Delta G^\circ = 2.303RT \log K_e$$

If both ΔH and ΔS are negative, the reaction will be spontaneous

A. at high T

B. at low T

C. at all T

D. at absolute zero

Answer: B

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228. Match the Column-I with Column-II

<u>Column-I</u>	<u>Column-II</u>
(A) A process carried out infinitesimally slowly	(P) Adiabatic
(B) A process in which no heat enters or leaves the system	(Q) $\Delta E = 0$
(C) A process carried out at constant temperature	(R) Reversible
(D) Cyclic process	(S) Isothermal
(E) Isochoric process	(T) $q_v = \Delta E$

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229. Match the Column-I with Column-II

Column-I

Column-II

- (A) $\text{H}_2(\text{g}) + \text{Cl}_2(\text{g}) \rightleftharpoons 2\text{HCl}(\text{g})$ (P) $\Delta H = \Delta U + RT$
(B) $\text{N}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{NO}(\text{g})$ (Q) $\Delta H = \Delta U$
(C) $\text{H}_2(\text{g}) + \text{I}_2(\text{s}) \rightleftharpoons 2\text{HI}(\text{g})$ (R) $\Delta H = \Delta U - 2RT$
(D) $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$ (S) $\Delta H < \Delta U$
(E) Perfectly crystalline solid (T) $\lim_{T \rightarrow 0} S \rightarrow 0$

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230. Match the Column-I with Column-II

Column-I

Column-II

- (A) $\Delta S_{\text{Total}} > 0$ (P) W_{useful} done by system
(B) $\Delta S_{\text{Total}} < 0$ (Q) Second law of thermodynamics
(C) ΔH (R) Non-spontaneous
(D) $-\Delta G$ (S) $\Delta U + \Delta nRT$

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231. Match the Column-I with Column-II

Column-I

Column-II

(A) $\frac{\Delta H_2 - \Delta H_1}{T_2 - T_1}$

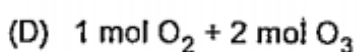
(P) Trouton equation

(B) $\frac{\Delta U_2 - \Delta U_1}{T_2 - T_1}$

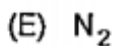
(Q) Effect of temperature on the heat of reaction

(C) $\frac{\Delta H_{\text{fusion}}}{T_{\text{mp}}} = \Delta S_{\text{fusion}}$

(R) Kirchoff's equation



(S) $C_P = \frac{7}{2}R$



(T) $C_P = \frac{23}{6}R$



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232. The enthalpy of neutralisation of oxalic acid by a strong base is -25.4 kcal/mol. The enthalpy of neutralisation of strong acid and strong base is $-13.7 \text{ kcal eq}^{-1}$. What is the enthalpy of dissociation (in kcal mol⁻¹) of oxalic acid ?



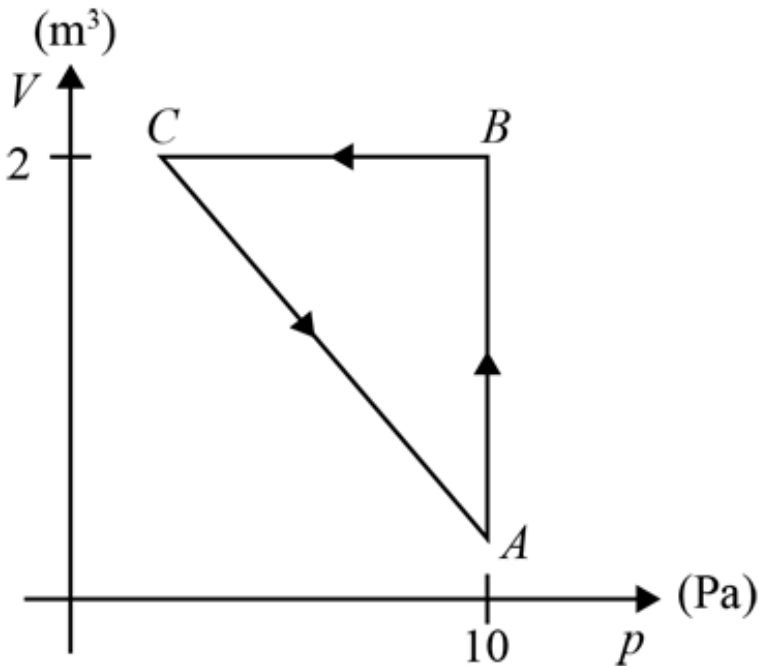
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233. 4.48 L of an ideal gas at STP requires 12 calories to raise its temperature by 15°C at constant volume. What is the C_p (in cal) of the gas ?



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234. An ideal gas is taken through the cycle $A \rightarrow B \rightarrow C \rightarrow A$ as shown in figure. If net heat supplied to the gas in the cycle is $+5\text{ J}$, what is the work done (in J) by the gas in the process $C \rightarrow A$?



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235. One mole of an ideal gas at 300 K is expanded ' isothermally from an Initial volume of (litre to 10 litre. What is the value of ΔE for this process ?

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236. 1 mole of an ideal gas is allowed to expand isothermally at $27^\circ C$ until its volume is tripled. What is the value of ΔS of universe when the expansion is carried out reversibly ?

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237. 1 mol of an ideal gas undergoes reversible isothermal expansion from an initial volume V_1 to final volume $10 V_1$ and does 10 kJ of work. The initial pressure was 1×10^7 Pa, Calculate the value of V_1 .

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238. 1 mole of an ideal gas undergoes reversible isothermal expansion from an initial volume V_1 to a final volume $10V_1$, and does 10 kJ of work. The initial pressure was 1×10^7 Pa.

If there were 2 mole of gas, what must its temperature would have been ?

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239. Two litre of N_2 at $0^\circ C$ and 5 atm pressure are expanded isothermally against a constant external pressure of 1 atm until the pressure of gas reaches 1 atm. Assuming gas to be ideal, calculate work of expansion.

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240. A 1.0 mol of ideal gas, initially at 10 atm and 300 K is allowed to expand isothermally to 1.0 atm. as follows :

Inversible expansion against 1.0 atm

Determine the net work done by the system and describe the whole processes on p-V diagram.

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241. A 1.0 mol of ideal gas, initially at 10 atm and 300 K is allowed to expand isothermally to 1.0 atm. as follows :

Inversible expansion against 1.0 atm

Determine the net work done by the system and describe the whole processes on p-V diagram.

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242. A sample of argon gas at 1 atm pressure and $27^{\circ}C$ expands reversibly and adiabatically from $1.25dm^3$ to $2.50dm^3$, Calculate the enthalpy change in this process, $C_{v,m}$ for argon is $12.48JK^{-1}mol^{-1}$,

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243. One mole of diatomic ideal gas undergoes a cyclic process ABC as shown in figure. The process BC is adiabatic. The temperatures at A, B and C are 400K, 800K and 600K respectively. Choose the correct statement:

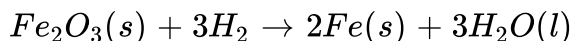
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- A. The Change in the Internal energy in the process AB is $-350R$
- B. The Change in the Internal energy in the process BC is $-500R$
- C. The Change in the Internal energy in the whole cyclic process is $250R$
- D. The Change in the Internal energy in the process CA is $700R$

Answer:

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244. Calculate, the ΔH at 85° for the reaction :

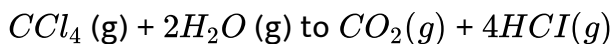


The data are : $\Delta(298)^\circ = -33.29 \text{ kJ/mol}$ and

Substance	$\text{Fe}_2\text{O}_3(\text{s})$	$\text{Fe}(\text{s})$	$\text{H}_2\text{O}(\text{l})$	$\text{H}_2(\text{g})$
$C_p^\circ \text{ J/K mol}$	103.8	25.1	75.3	28.8

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245. The standard heats of formation at 298 K for $\text{CCl}_4(\text{g})$, $\text{H}_2\text{O}(\text{g})$, $\text{CO}_2(\text{g})$ and $\text{HCl}(\text{g})$ are -25.5 , -57.8 , -94.1 and $-22.1 \text{ kcal mol}^{-1}$ respectively, calculate $\Delta H_{298\text{K}}^\circ$, for the reaction :



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246. Show that the reaction $\text{CO}(\text{g}) + \frac{1}{2}\text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g})$ at 300 K is spontaneous and exothermic, when the standard entropy change is $-0.094 \text{ kJ mol}^{-1} \text{ K}^{-1}$. The standard Gibbs free energies of formation for CO_2 and CO are -394.4 and -137 kJ mol^{-1} , respectively.

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247. An isolated system is that system in which

- A. there is no exchange of energy with the surroundings
- B. there is exchange of mass and energy with the surroundings
- C. there is no exchange of energy and mass with the surroundings
- D. there is exchange of mass with the surroundings

Answer: C



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248. For a cycle process the condition(s) is/are

- A. $\Delta U = 0$
- B. $\Delta H = 0$
- C. $\Delta U > 0$ and $\Delta H > 0$
- D. Both $\Delta U = 0$ and $\Delta H = 0$

Answer: D

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249. Identify the intensive property from the following

- A. enthalpy and temperature
- B. volume and temperature
- C. enthalpy and volume
- D. temperature and refractive index

Answer: D

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250. The mathematical form of the first law of thermodynamics when q is the heat supplied and w the work done by the system is

A. $\Delta U = q + w$

B. $\Delta U = q - w$

C. $\Delta U = w - q$

D. $\Delta U = -q - w$

Answer: B

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251. Which one of the following is a state function?

A. Heat

B. Work

C. Internal energy

D. Potential energy

Answer: C,D

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252. Calculate the maximum work done by an ideal gas when pressure on 10 g of H_2 is reduced from 20 to 2 atm at a constant temperature 300 K.

A. -28.72 kJ

B. -28.72 J

C. -2.872 kJ

D. -2.872 J

Answer: A



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253. The work done when one mole of an ideal gas expands irreversibly under adiabatic condition can be given by the expression is

A. $-P_{ext}(V_2 - V_1)$

B. $-2.303RT \log\left(\frac{V_2}{V_1}\right)$

C. $-2.303RT \log\left(\frac{P_2}{P_1}\right)$

D. γ

Answer: A



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254. Two mole of an ideal gas is expanded isothermally and reversibly from 1 litre to 10 litre at 300 K. The enthalpy change (in kJ) for the process is

A. 11.4 kJ

B. -11.4 kJ

C. 0 kJ

D. 4.8 kJ

Answer: C



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255. When a monoatomic gas undergoes an adiabatic process, its temperature and volume are related by the equation $TV^n = \text{constant}$, the value of n will be

- A. 1.33
- B. 0.66
- C. 2.33
- D. 1

Answer: B

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256. A monoatomic gas X and a diatomic gas Y, both initially at the same temperature and pressure are compressed adiabatically (separately) from a volume V to $V/2$, which gas will be at higher temperature?

- A. X

B. Y

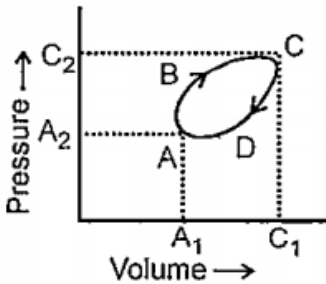
C. both are same

D. can't say

Answer: A

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257. A thermodynamic system undergoes a cyclic change as represented in the following P-V diagram



The net work done during the complete cycle is given by the area

A. Cycle ABCDA

B. AA_1C_1CDA

C. AA_2C_2C

D. half of area bounded by the curve

Answer: A

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258. If w_1, w_2, w_3 and w_4 are work done in isothermal, adiabatic, isobaric and isochoric reversible processes, then the correct sequence (for expansion) would be

A. $w_1 > w_2 > w_3 > w_4$

B. $w_3 > w_1 > w_2 > w_4$

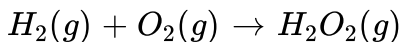
C. $w_3 > w_2 > w_1 > w_4$

D. $w_3 > w_2 > w_4 > w_1$

Answer: B

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259. Calculate ΔH for this reaction,



$$B_{H-H} = 436kJ, B_{O-O} = 142kJ, B_{O=O} = 499kJ, B_{O-H} = 460kJ$$

A. -127 kJ

B. -209 kJ

C. -484 kJ

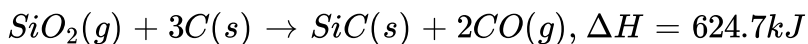
D. -841 kJ

Answer: A



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260. How much heat is absorbed in complete reaction of 3.00 g of SiO_2 with excess of carbon in the reaction given below?



A. $1.33 \times 10^4 kJ$

B. $1.13 \times 10^5 \text{ kJ}$

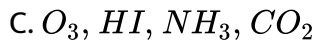
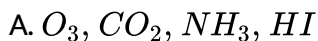
C. 5.06 kJ

D. 31.2 kJ

Answer: D

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261. Standard enthalpies of formation of O_3 , CO_2 , NH_3 and HI are 142.2, -393.2, -46.2 and $+25.9 \text{ kJ mol}^{-1}$ respectively. The order of their increasing stabilities will be



Answer: C



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262. What should be the amount of heat released when an aqueous solution containing 0.5 mole of NH_4^+ is mixed with 0.3 mole of OH^- ? (enthalpy of neutralisation is -57.1 kJ)

- A. 28.5 kJ
- B. 17.1 kJ
- C. 45.7 kJ
- D. 1.7 kJ

Answer: B



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263. The difference between the heats of reaction at constant pressure and a constant volume for the reaction $2C_6H_6(l) + 15O_2(g) \rightarrow 12CO_2(g) + 6H_2O(l)$ at $25^\circ C$ in kJ is

- A. -7.43
- B. +3.72
- C. -3.72
- D. +7.43

Answer: A

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264. The value of ΔH_{comb} of four gases C_2H_6 , C_2H_4 , C_2H_{10} , CH_4 are given to be -1560, -1411, -2877 and $-890.4 kJ mol^{-1}$ respectively. Which of these is the best fuel?

- A. C_2H_6
- B. C_2H_4
- C. C_2H_{10}
- D. CH_4

Answer: D



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265. A solution of 500 mL of 0.2 M KOH and 500 mL of 0.2 M HCl are mixed and stirred, the rise in temperature is T_1 . The experiment is repeated using 250 mL of each of the solution, the temperature rise is T_2 . Which of the following is true?

A. $T_1 = T_2$

B. $T_1 = 2T_2$

C. $T_1 = 4T_2$

D. $T_2 = 9T_1$

Answer: A



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266. The standard heat of formation of $NO_2(g)$ and $N_2O_4(g)$ are 8.0 and $2.0 \text{ kcal mol}^{-1}$ respectively. The heat of dimerization of NO_2 in kcal is:

- A. 10
- B. -6
- C. -12
- D. -14

Answer: D



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267. Given enthalpy of formation of $CO_2(g)$ and $CaO(s)$ are -94.0 kJ and -152 kJ respectively and the enthalpy of the reaction,

$CaCO_3(s) \rightarrow CaO(s) + CO_2(g)$ is 42 kJ. The enthalpy of formation of $CaCO_3(s)$ is

- A. -42 kJ

B. -202 kJ

C. +202 kJ

D. -288 kJ

Answer: D

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268. $H_2(g) \rightarrow 2H(g)$, $\Delta H = 104.2 \text{ kcal}$. The bond energy of H-H bond is

A. 104.2 kcal

B. 208.4 kcal

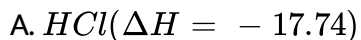
C. 52.1 kcal

D. zero

Answer: A

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269. Which compound will absorb the maximum amount of heat when dissolved in the same amount of water? (Integral heats of solution at $25^{\circ}C$ in kcal/mol of each solute are given in brackets)



Answer: C

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270. Heat evolved in the reaction, $H_2 + Cl_2 \rightarrow 2HCl$ is 182 kJ. Bond energies of H-H and Cl-Cl are 430 and 242 kJ/mol respectively. The H-Cl bond energy is



B. 427kJmol^{-1}

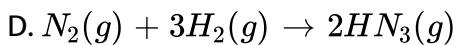
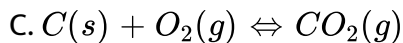
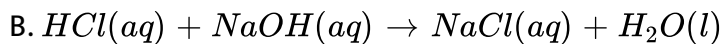
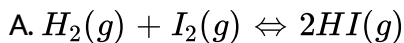
C. 336kJmol^{-1}

D. 154kJmol^{-1}

Answer: B

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271. For which change $\Delta H \neq \Delta U$?



Answer: D

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272. The enthalpies of combustion of carbon and carbon monoxide are -393.5 and -283kJmol^{-1} respectively. The enthalpy of formation of carbon monoxide per mole is

- A. -110.5 kJ
- B. 676.5 kJ
- C. -676.5 kJ
- D. 110.5 kJ

Answer: A

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273. A particular reaction has $\Delta H = +250\text{kJmol}^{-1}$, which of the following statements is/are true?

- I. Heat is given to the surroundings.
- II. The reaction is endothermic.
- III. Products are more stable than the reactants.

A. all are true

B. only II and III are true

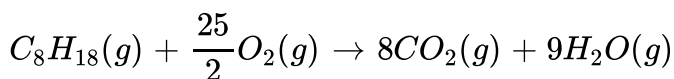
C. only I, III are true

D. only II is true

Answer: D

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274. Combustion of octane takes place in an automobile engine. The homogeneous equation of combustion is:



The signs of ΔH , ΔS and ΔG for the reaction will be

A. +ve, -ve, +ve

B. -ve, +ve, -ve

C. -ve, +ve, +ve

D. +ve, +ve, -ve

Answer: B

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275. Estimate the minimum temperature above which the reaction will be spontaneous when $\Delta H = -176\text{kJ}$ and $\Delta S = -284.5\text{JK}^{-1}$

A. 582 K

B. 834 K

C. 634 K

D. 619 K

Answer: D

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276. ΔH and ΔS for a reaction are $+30.558\text{kJmol}^{-1}$ and $0.066\text{kJK}^{-1}\text{mol}^{-1}$ at 1 atm pressure. The temperature at which free

energy change is zero and the nature of the reaction below this temperature is

- A. 483 K, spontaneous
- B. 443 K, non-spontaneous
- C. 443 K, spontaneous
- D. 463 K, non-spontaneous

Answer: D



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277. For which of these processes is the value of ΔS negative

- I. Sugar is dissolved in water.
- II. Steam condenses on a surface
- III. $CaCO_3$ is decomposed into CaO and CO_2

- A. I only
- B. II only

C. I and II only

D. II and III only

Answer: B



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278. The direct conversion of A to B is difficult, hence it is carried out by the following shown path:

$A \rightarrow C \rightarrow D \rightarrow B$, Given $\Delta S_{A \rightarrow C} = 50unit$,

$\Delta S_{C \rightarrow D} = 30unit$ and $\Delta S_{B \rightarrow D} = 20unit$,

Therefore, total entropy change $\Delta S_{A \rightarrow B}$ is

A. $+100unit$

B. $+60unit$

C. $-100unit$

D. $-60unit$

Answer: B



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279. Identify the correct statement regarding a spontaneous process

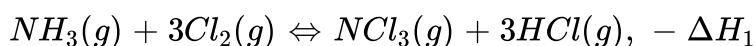
- A. For a spontaneous process in an isolated system, the change in entropy is positive
- B. Endothermic processes are never spontaneous
- C. Exothermic process are always spontaneous
- D. Lowering of energy in the reaction process is the only criterion for spontaneity

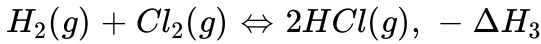
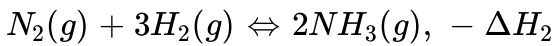
Answer: A



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280. Given,





The heat of formation of $NCl_3(g)$ in terms of ΔH_1 , ΔH_2 and ΔH_3 is

A. $\Delta H_f = \Delta H_1 + \frac{\Delta H_2}{2} - \frac{3}{2}\Delta H_3$

B. $\Delta H_f = -\Delta H_1 + \frac{\Delta H_2}{2} - \frac{3}{2}\Delta H_3$

C. $\Delta H_f = -\Delta H_1 - \frac{\Delta H_2}{2} - \frac{3}{2}\Delta H_3$

D. None of the above

Answer: D



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281. C_V value of He is always $3/2 R$. C_V value for hydrogen at low temperature is also $3R/2$. C_V value of hydrogen gas at moderate temperature is

A. $3/2 R$

B. $5/2 R$

C. $> \frac{5}{2}R$

D. none

Answer: B



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282. An ideal gas is allowed to expand both reversibly and irreversibly in an isolated system. If T_i is the initial temperature and T_f is the final temperature, which of the following statements is correct?

A. $(T_f)_{irr.} > (T_f)_{rev.}$

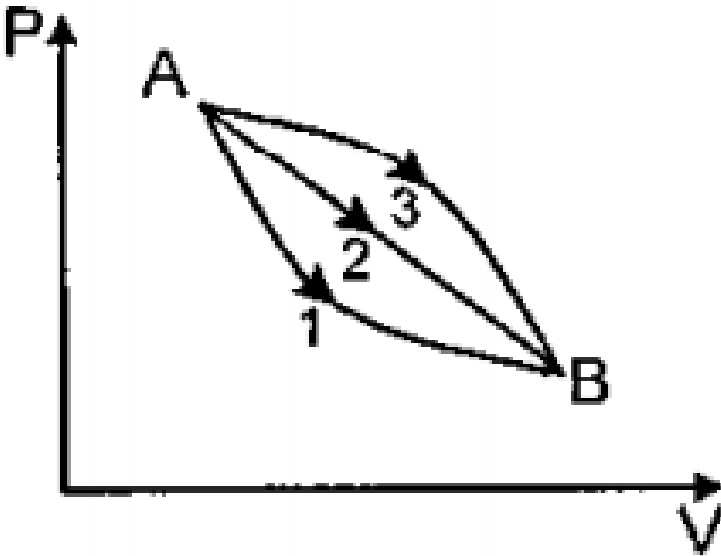
B. $T_f > T_i$ for reversible process and $T_f = T_i$ for irreversible process

C. $(T_f)_{rev.} > (T_f)_{irr.}$

D. $T_f = T_i$ for both reversible and irreversible process

Answer: A

283. A given mass of gas expands from the state A to state B by three paths 1, 2 and 3 as shown in the figure. If w_1 , w_2 and w_3 respectively be the work done by the gas along three paths then



A. $w_1 > w_2 > w_3$

B. $w_1 < w_2 < w_3$

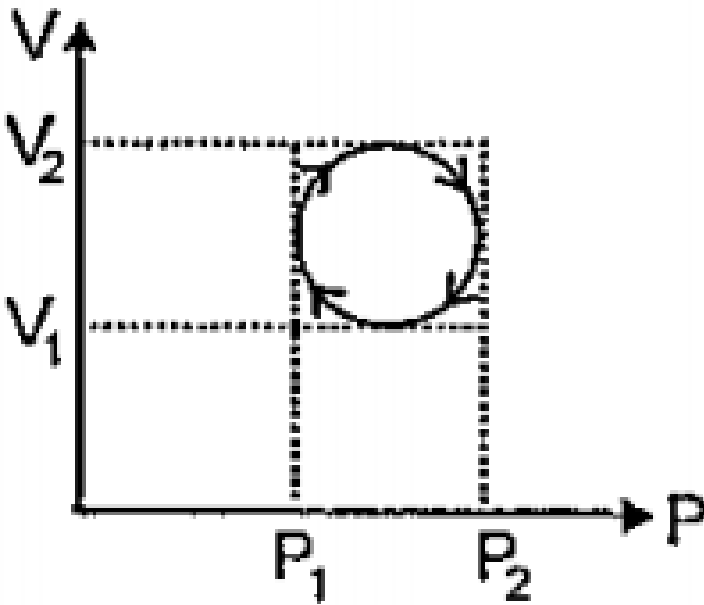
C. $w_1 = w_2 = w_3$

D. $w_2 < w_3 < w_1$

Answer: B

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284. In the cyclic process shown in P-V diagram, the magnitude of the work done is:



A. $\pi \left(\frac{P_2 - P_1}{2} \right)$

B. $\pi \left(\frac{V_2 - V_1}{2} \right)$

C. $\frac{\pi}{4}(P_2 - P_1)(V_2 - V_1)$

D. $\pi(V_2 - V_1)$

Answer: C

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285. The heat liberated when 1.89 g of benzoic acid is burnt in a bomb calorimeter at $25^\circ C$ increase the temperature of 18.94 kg of water by $0.632^\circ C$. If the specific heat of water at $25^\circ C$ is 0.998 cal/g-deg, the value of the heat of combustion of benzoic acid is

A. 881.1 kcal

B. 771.4 kcal

C. 981.1 kcal

D. 871.2 kcal

Answer: B

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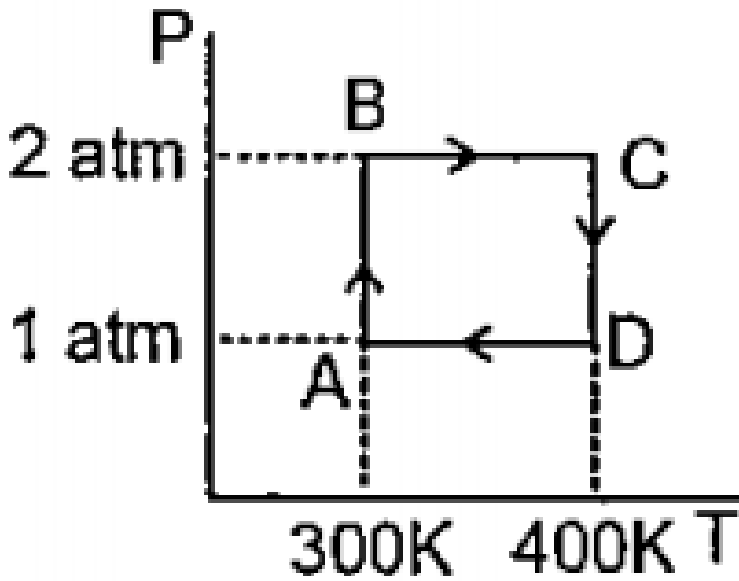
286. 1 mole of NH_3 gas at $27^\circ C$ is expanded in reversible adiabatic condition to make volume 8 times ($\gamma = 1.33$). Final temperature and work done respectively are

- A. 150 K, 900 cal
- B. 150 K, 400 cal
- C. 250 K, 1000 cal
- D. 200 K, 800 cal

Answer: A

287. Two moles of Helium gas undergoes a cyclic process as shown in figure. Assuming gas to be ideal, what is the net work involved in the

cyclic process?



A. $-138R \ln 2$

B. $+150R \ln 2$

C. $+200R \ln 2$

D. $-200R \ln 2$

Answer: A



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288. Under the same conditions how many mL of 1 M KOH and 0.5M H_2SO_4 solution, respectively when mixed for a total volume of 100 mL produce the highest rise in temperature?

A. 67 and 33

B. 33 and 67

C. 40 and 60

D. 50 and 50

Answer: D



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289. Heat of neutralisation of oxalic acid is $-106.7kJmol^{-1}$ using NaOH.

Hence ΔH of $H_2C_2O_4 \Leftrightarrow C_2O_4^{2-} + 2H^+$ is

A. 5.88 kJ

B. -5.88 kJ

C. -13.7 kJ

D. 7.5 kJ

Answer: D



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290. If a and b are arbitrary extensive variables then

A. $(a+b)$ is an extensive variable

B. $(a+b)$ is an intensive variable

C. a/b is an intensive variable

D. da/db is an intensive variable

Answer:



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291. During an adiabatic reversible expansion of an ideal gas

- A. internal energy of the system decreases
- B. temperature of the system decreases
- C. The value of γ changes
- D. pressure increases

Answer:



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292. Which is/are irreversible process(es)?

- A. Mixing of two gases by diffusion
- B. Evaporation of water at 373 K and 1 atm in a closed system
- C. Dissolution of NaCl in water
- D.

Answer:

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293. Which of the following is/are thermodynamically stable?

A. C (diamond)

B. C (graphite)

C. P_4 (white)

D. P_4 (black)

Answer:

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294. Which of the following expressions represent(s) work done by the isothermal reversible expansion of the ideal gas?

- A. $-2.303nRT \log\left(\frac{V_2}{V_1}\right)$
- B. $-2.303nRT \log\left(\frac{P_1}{P_2}\right)$
- C. $-2.303nRT \log\left(\frac{V_1}{V_2}\right)$
- D. $-2.303nRT \frac{dP}{dT} = \frac{\Delta V}{\Delta S}$

Answer:

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295. For the adiabatic expansion of an ideal gas

- A. $PV^\gamma = \text{constant}$
- B. $TV^{\gamma-1} = \text{constant}$
- C. $TP^{1-\gamma} = \text{constant}$
- D. none of the above

Answer:

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296. Which of the following is/are correct?

- A. $\Delta H = \Delta U + \Delta(PV)$ when P and V both change
- B. $\Delta H = \Delta U + P\Delta V$ When pressure is constant
- C. $\Delta H = \Delta U + V\Delta P$ When volume is constant
- D. $\Delta H = \Delta U + P\Delta V + V\Delta P$ when P and V both change

Answer:



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297. Which of the following statement(s) is/are correct?

- A. The reaction between strong acid and strong base takes place with evolution of energy.

B. $|\Delta H_N|$ of weak acid/strong base is less than $|\Delta H_N|$ of strong acid/strong base.

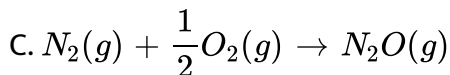
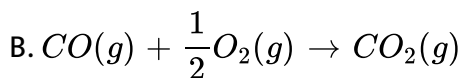
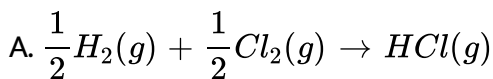
C. ΔH_N of strong acid/strong base is equal to the ΔH of formation of $H_2O(l)$ from its ions in aqueous medium.

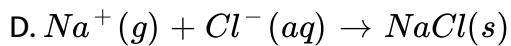
D. ΔH_{net} (weak acid/strong base) - ΔH_{net} (strong acid/strong base) = $\Delta H_{\text{ionization}}$ (weak acid)

Answer:

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298. Which of the following reactions do(es)n't represent the standard state enthalpy of formation reaction?





Answer:

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299. In a reaction ΔH and ΔS both are more than zero. In which of the following cases, the reaction would not be spontaneous?

A. $\Delta H > T\Delta S$

B. $T\Delta S > \Delta H$

C. $\Delta H = T\Delta S$

D. $\Delta G < 0$

Answer:

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300. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : Heat and work are "definite quantities".

Statements-II : Heat and work are not properties of a system, their values depend on the path of the process and vary accordingly.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: D



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301. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : There is no change in internal energy for an ideal gas at constant temperature.

Statement-II : Internal energy of an ideal gas is a function of temperature only.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: A



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302. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : The heat absorbed during the isothermal expansion of an ideal gas against vacuum is zero.

Statement-II : The volume occupied by the molecules of an ideal gas is zero.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: B



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303. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : The magnitude of the work involved in an isothermal expansion is greater than that involved in an adiabatic expansion.

Statement-II : P-V curve (P on y-axis and V on x-axis) decrease more rapidly for reversible adiabatic expansion compared to reversible isothermal expansion starting from same initial state.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: A



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304. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : Entropy change in reversible adiabatic expansion of an ideal is zero.

Statement-II : The increase in entropy due to volume increase just compensate the decrease in entropy due to fall in temperature.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: A



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305. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : The standard free energy changes of all spontaneously occurring reactions are negative.

Statement-II : The standard free energies of the elements in their standard states at 1 bar and 298 K are taken as zero.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: B



306. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : Enthalpy and entropy of any elementary substance in the standard states are taken as zero.

Statement-II : At absolute zero, particles of the perfectly crystalline substance become completely motionless.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: D



307. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : A reaction which is spontaneous and accompanied by decrease of randomness must be exothermic.

Statement-II : All exothermic reactions are accompanied by decrease of randomness.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: C



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308. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : $|\Delta_f H|$ of $H_2O(l) > |\Delta_f H|$ of $H_2O(g)$

Statement-II : $\Delta H_{condensation}$ is negative.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: A



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309. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : All combustion reactions are exothermic.

Statement-II : Enthalpies of products are greater than enthalpies of reactants $\left(\sum V_P \Delta_f H(P) > \sum V_R \Delta_f H(R) \right)$

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: C



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310. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : Enthalpy of neutralization of CH_3COOH by NaOH is less than that of HCl by NaOH.

Statement-II : Enthalpy of neutralization of CH_3COOH is less because of the absorption of heat in the ionization process.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: A



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311. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : Internal energy of a real gas may change during expansion at constant temperature.

Statement-II : Internal energy of a real gas is a function of temperature and volume.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: A



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312. This question has Statement I and Statement II. Of the four choices given after the Statements, choose the one that best describes the two Statements.

Statement-I : Work is a state function which is expressed in joule.

Statement-II : Work appears only at the boundary of the system.

- A. Statement-I is true, Statement-II is true, Statement-II is a correct explanation of Statement-I
- B. Statement-I is true, Statement-II is true, Statement-II is not a correct explanation of Statement-I
- C. Statement-I is true, Statement-II is false.
- D. Statement-I is false, Statement-II is true.

Answer: D



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313.

C_P for monoatomic gas is

A. $5R/2$

B. $3R/2$

C. $7R/2$

D. R

Answer: A



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314. Molar heat capacity is heat required to raise the temperature of one mole of material by one degree, since heat is not a state function, the amount of heat required to produce a given change in its state depends on the path followed.

$C_P = \text{specific heat} \times \text{Molecular weight}$. It is measured at constant pressure

$C_V = \text{specific heat} \times \text{Molecular weight}$. It is measured at constant volume

Answer the following questions based on above passage:

The specific heat of a gas at constant volume is 0.075 cal/g. Predict the atomicity of the gas. Molar mass of gas is 40 g mol^{-1}

A. 1

B. 2

C. 3

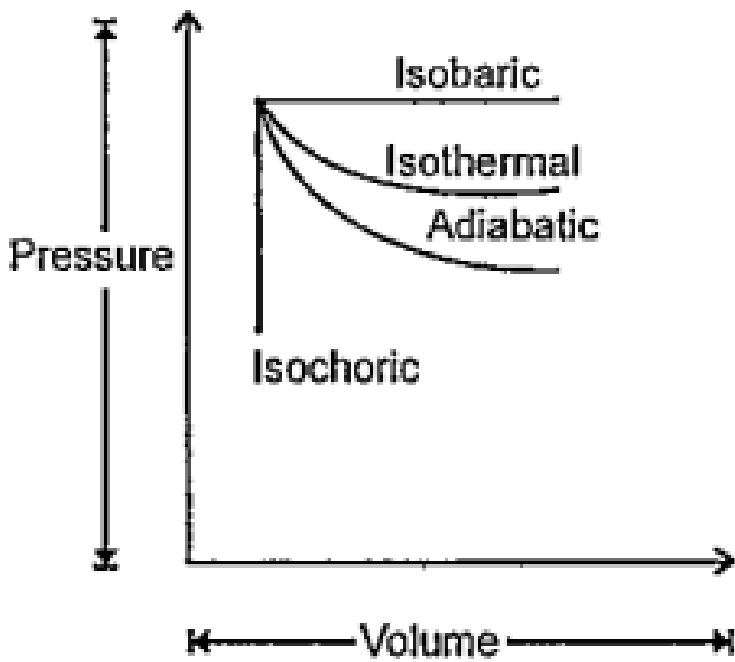
D. none of the above

Answer: A



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315. Observe the following graphic representation of four basic thermodynamic processes.



Answer the following questions based on above passage:

Which of the following is true isochoric process?

- A. $\Delta V = 0$
- B. $\Delta E = dq + P\Delta V$
- C. $\Delta E = q_v$
- D. All of these

Answer: D



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316. Match Column-I with Column-II

Column-I

- (A) Reversible isothermal expansion of an ideal gas
- (B) Reversible adiabatic compression
- (C) Irreversible adiabatic expansion of an ideal gas
- (D) Irreversible isothermal compression of an ideal gas

Column-II

- (P) $w = -2.303nRT \log \left(\frac{V_2}{V_1} \right)$
- (Q) $PV^\gamma = \text{constant}$ of an ideal gas
- (R) $w = \frac{nR}{(\gamma-1)} (T_2 - T_1)$
- (S) $\Delta H = 0$



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317. Match Column-I with Column-II

Column-I

Column-II

- | | |
|-----------------|-------------------------|
| (A) Enthalpy | (P) Intrinsic property |
| (B) Temperature | (Q) Path function |
| (C) Free energy | (R) Function of E, P, V |
| (D) Work | (S) State function |

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318. 2 g of an organic compound is burnt completely in a bomb calorimeter at 289 K. The final temperature of calorimeter was found to be 298.5 K. Calorimeter constant is $400JK^{-1}$ and molar mass of the compound is 50 g. What is the enthalpy of combustion per mol of the compound in kJ unit?

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319. When a current of x ampere from a 20.9 V battery is used for 4.0 s to heat 100 g water, its temperature increased by 1°C . Heat capacity of water is 4.18 JK^{-1} . If all the electrical energy is converted into thermal energy, what is x ?

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320. When 5.0 ml of 0.1 M HCl solution is mixed with 5.0 ml of a 0.1 M NaOH solution, the temperature of solution increases by $2\text{ }^\circ\text{C}$. Which of the following can be predicted accurately from this observation?

- A. If 10 ml of same HCl is mixed with 10 ml of same NaOH , the temperature will be 4°C
- B. If 10 ml of 0.5 M HCl is mixed with 10 ml of 0.5 M NaOH , the temperature will be 2°C
- C. If 5 ml of 0.1 M HCl is mixed with 5 ml , 0.1 M NH_3 solution, the temperature rise will be less than $2\text{ }^\circ\text{C}$

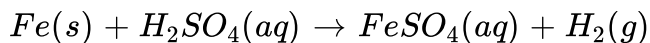
D. If 5 ml, 0.1 M CH₃COOH solution is mixed with 5 ml, 0.1 M NaOH

solution, the temperature rise will be less than 2 °C

Answer:

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321. Calculate the work done by the reaction (in L · atm)



when 0.3125 mole of H₂ gas is collected at 273K at 1.0 atm.

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322. Calculate entropy change when 10 moles of an ideal gas expands reversibly and isothermally from an initial volume of 10 litre to 100 litre at 300K.

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323. A gas present in a cylinder filled with a frictionless piston expands against a constant pressure of 1 atm from a volume of 2 litre to a volume of 6 litre. In doing so, it absorbs 800 J heat from surroundings. Determine increase in internal energy of process.

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324. Work done in expansion of an ideal gas from 4 litre to 6 litre against a constant external pressure of 2.5 atm was used to heat up 1 mole of water at 293K. If specific heat of water is $4.184 \text{ J g}^{-1} \text{ K}^{-1}$, what is the final temperature of water?

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325. Calculate the maximum work done in expanding 16 g of oxygen at 300K and occupying a volume of 5 dm^3 isothermally until the volume becomes 25 dm^3 .

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326. One mole of an ideal monoatomic gas is heated at constant pressure from $0^{\circ}C$ to $100^{\circ}C$.

Calculate work done.

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327. One mole of an ideal gas is heated at constant pressure from $0^{\circ}C$ to $100^{\circ}C$. (a) Calculate work done. (b) If the gas is expanded isothermally and reversibly at $0^{\circ}C$ from 1 atm to some other pressure P_1 . what must be the final pressure if the maximum work is equal to the work involved in (a)?

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328. Calculate ΔH° for $2Al + Fe_2O_3 \rightarrow 2Fe + Al_2O_3$ given that standard heat enthalpy of Fe_2O_3 and Al_2O_3 are -196.5 and

$$-399.1 \text{ kcal mol}^{-1}$$

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329. The heat liberated on complete combustion of 7.8 g benzene is 327 kJ. Calculate heat of combustion measured at constant volume and at 27°C .

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330. Calculate heat of combustion of benzene at constant pressure at 27°C . ($R = 8.3 \text{ J mol}^{-1} \text{ K}^{-1}$)

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331. The enthalpy change involved in the oxidation of glucose is $-2880 \text{ kJ mol}^{-1}$. Twenty five percent of this energy is available for muscular work. If 100 kJ of muscular work is needed to walk one kilometre,

what is the maximum distance that a person will be able to walk after eating 120 g of glucose?

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332. The heat of formation of ethane is -20.3 kcal. Calculate the bond energy of C-C bond in ethane if the heats of atomisation of carbon and hydrogen are respectively 170.9 and 52.1 kcal per mole and bond energy of C-H bond is 99.0 kcal.

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333. Calculate the resonance energy of N_2O from the following data

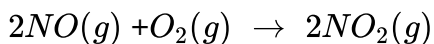
$\Delta_1 H^0$ of $N_2O = 82 \text{ kJ mol}^{-1}$ for the reaction

$N_2(g) + \frac{1}{2}O_2(g) \rightarrow N_2O(g)$ Bond energy of $N = N$, $N = N$, $O = O$

and $N = O$ bonds are $946, 418, 498$ and 607 kJ mol^{-1} respectively.

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334. The following reaction is performed at 298 K.



The standard free energy of formation of $NO(g)$ is 86.6 kJ/mol at 298 K.

What is the standard free energy of formation of $NO_2(g)$ at 298 K? ($K_p =$

$$1.6 \times 10^{12}$$
)

A. $86600 + R(298)\ln(1.6 \times 10^{12})$

B. $86600 - \left(\frac{\ln(1.6 \times 10^{12})}{R(298)} \right)$

C. $0.5[2 \times 86,600 - R(298)\ln(1.6 \times 10^{12})]$

D. $R(298)\ln(1.6 \times 10^{12}) - 86600$

Answer: C



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335. The standard Gibbs energy change at 300 K for the reaction

$2A \rightleftharpoons B + C$ is 2494.2 J. At a given time, the composition of the reaction

mixture is $[A] = 1/2$, $[B] = 2$ and $[C] = 1/2$. The reaction proceeds in the: $[R = 8.314 \text{ J/K/mol}, e = 2.718]$

- A. reverse direction because $Q > K_c$
- B. forward direction because $Q < K_c$
- C. reverse direction because $Q < K_c$
- D. forward direction because $Q > K_c$

Answer: A



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336. For the reaction $X_2Y_4(l) \rightarrow 2XY_2(g)$ at 300 K the values of ΔU and ΔS are 2 kCal and 20 Cal K^{-1} respectively. The value of ΔG for the reaction is

- A. -3400 cal
- B. 3400 cal
- C. -2800 cal

D. 2000 cal

Answer: C

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337. The values of ΔH and ΔS of a certain reaction are -400kJmol^{-1} and -20kJmol^{-1} respectively. The temperature below which the reaction is spontaneous is

A. 100°K

B. 20°C

C. 20°K

D. 120°C

Answer: C

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338. The enthalpy of vaporization of a certain liquid at its boiling point of $35^{\circ}C$ is $24.64kJmol^{-1}$. The value of change in entropy for the process is

A. $704JK^{-1}mol^{-1}$

B. $80JK^{-1}mol^{-1}$

C. $24.64JK^{-1}mol^{-1}$

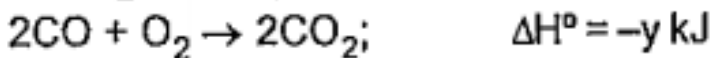
D. $7.04JK^{-1}mol^{-1}$

Answer: B



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339. Given that



The heat of formation of carbon monoxide will be

A. $(y-2x)/2$

B. $y+2x$

C. $2x-y$

D. $(2x-y)/2$

Answer: A

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340. The value of ΔH for cooling 2 mole of an ideal monoatomic gas from $225^\circ C$ to $125^\circ C$ at constant pressure will be $C_P = \frac{5}{2}R$

A. $250 R$

B. $-500 R$

C. $500 R$

D. $-250 R$

Answer: B

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341. For a spontaneous process, the correct statement(s) is (are)

A. $(\Delta G_{system})_{T,P} > 0$

B. $(\Delta S_{system}) + (\Delta S_{surroundings}) > 0$

C. $(\Delta G_{system})_{T,P} < 0$

D. $(\Delta U_{system})_{T,V} > 0$

Answer: B::C



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342. Which of the following statements is correct for the spontaneous absorption of a gas?

A. ΔS is negative and therefore, ΔH should be highly positive

B. ΔS is negative and therefore, ΔH should be highly negative

C. ΔS is negative and therefore, ΔH should be negative

D. ΔS is positive are therefore, ΔH should also be highly positive

Answer: B

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343. For the reaction $X_2O_4(l) \rightarrow 2XO_2(g)$, $\Delta U = 2.1 \text{ kcal}$, $\Delta S = 20 \text{ cal } K^{-1}$ at 300k

Hence, ΔG is

- A. 2.7 kcal
- B. -2.7 kca,
- C. 9.3 kcal
- D. -9.3 kcal

Answer: B

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344. The ratio of heats liberated at 298K from the combustion of one kg of coke and byburning water gas obtained from 1kg of coke is (Assume coke to be 100 % carbon). (Given : enthalpies of combustion of CO_2 , CO and H_2 are 393.5 kJ, 285 kJ, 285 kJ respectively all at 298K)

A. 0.79:1

B. 0.69:1

C. 0.80:1

D. 0.96:1

Answer: B



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345. Conversion of oxygen into ozone is non-spontaneous at

A. all temperature

B. high temperature

C. room temperature

D. low temperature

Answer: A



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346. For an ideal binary liquid mixture

A. $\Delta S_{mix} = 0, \Delta G_{mix} = 0$

B. $\Delta H_{mix} = 0, \Delta S_{mix} < 0$

C. $\Delta V_{mix} = 0, \Delta G_{mix} > 0$

D. $\Delta S_{mix} > 0, \Delta G_{mix} < 0$

Answer: D



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347. The work done when one mole of an ideal gas is compressed from a volume of 5m^3 to 1dm^3 at 300 K, under a pressure of 100 kPa is

- A. 499.8 kJ
- B. -499.9 kJ
- C. -99.5 kJ
- D. 42495 kJ

Answer: A



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348. Assuming enthalpy of combustion of hydrogen at 273K is -286 kJ/mol and enthalpy of fusion of ice at the same temperature to be $+6.0\text{ kJ/mole}$, calculate enthalpy change during formation of 100g of ice.

- A. $+1622\text{kJ}$
- B. -1622 kJ

C. $+292kJ$

D. -292 kJ

Answer: B

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349. A piston filled with 0.04mole of an ideal gas expands reversibly from 50.0mL to 375mL at a constant temperature of 37.0°C . As it does so, it absorbs 208J of heat. The values of q and W for the process will be ($R = 8.314\text{J/mol K}$, $\ln 7.5 = 2.01$)

A. $q = +208J, W = -208J$

B. $q = -208\text{ J}, W = -208\text{ J}$

C. $q = -208J, W = +208J$

D. $q = +208J, W = +208J$

Answer: A

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350. If ΔG for a reaction is negative, you infer that the change is

- (1) spontaneous
- (2) non-spontaneous
- (3) reversible
- (4) irreversible
- (5) equilibrium

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351. In order to decompose 9g water 142.5 kJ heat is required. Hence, the enthalpy of formation of water is

- A. -142.5 kJ/mole
- B. +142.5 kJ/mole
- C. -285 kJ/mole
- D. +285 kJ/mole

Answer: C

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352. The enthalpy of combustion of $C_6H_6(l)$ is -3250 kJ. When 0.39 g of benzene is burnt in excess of oxygen in an open vessel, the amount of heat liberated is

- A. 16.25 J
- B. 16.25 kJ
- C. 32.5 J
- D. 32.5 J

Answer: B

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353. The intensive property among these quantities is

A. enthalpy

B. mass/volume

C. mass

D. volume

Answer: B



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354. Hess's law states that

A. the standard enthalpy of an overall reaction is the sum of the enthalpy changes in individual reactions

B. enthalpy of formation of a compound is same as the enthalpy of decomposition of the compound into constituent elements, but with opposite sign

C. at constant temperature the pressure of a gas is inversely proportional to its volume

D. the mass of a gas dissolved per litre of a solvent is proportional to the pressure of the gas in equilibrium with the solution

Answer: A

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355. A gas can expand from 100 mL to 250 mL. under a constant pressure of 2 atm. The work done by the gas is

A. 30.38 J

B. 25 J

C. 5 kJ

D. 16 J

Answer: A

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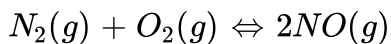
356. For a Process to be spontaneous

- A. ΔG must be -ve
- B. ΔG should be +ve
- C. ΔH must be -ve
- D. ΔS must be -ve

Answer: A

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357. Assertion ΔH and ΔE are almost the same for the reaction,



Reason All reactants and products are gases.

- A. If both Assertion and Reason are true and reason is correct explanation of Assertion
- B. If both Assertion and Reason are true but reason is not the correct explanation of Assertion
- C. If Assertion is true but Reason is false
- D. If both Assertion and Reason are false

Answer: B



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358. Assertion The increase in internal energy (ΔE) for the vaporisation of one mole of water at 1 atm and 373 K is zero.

Reason For all isothermal processes $\Delta E = 0$.

- A. if both Assertion and Reason are true and Reason is the correct explanation of Assertion

- B. if both Assertion and Reason are true but Reason is not the correct explanation of Assertion
- C. If Assertion is true but Reason is false
- D. If both Assertion and Reason are false

Answer: A

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359. Which of the following has ΔS° greater than zero?

- A. $CaO(s) + CO_2(g) \rightleftharpoons CaCO_3(s)$
- B. $NaCl(aq) \rightleftharpoons NaCl(s)$
- C. $NaNO_3(s) \rightleftharpoons Na^+(aq) + NO_3^-(aq)$
- D. $N_2(s) + 3H_2(g) \rightleftharpoons 2NH_3(g)$

Answer:

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360. The heat of reaction for

$C_{10}H_8(s) + 12O_2(g) \rightarrow 10CO_2(g) + 4H_2O(l)$ at constant volume is -1228.2 kcal at $25^\circ C$. The heat of reaction at constant pressure and same temperature is

- A. -1228.2 kcal
- B. -1229.3 kcal
- C. -1232.9 kcal
- D. -1242.6 kcal

Answer: C



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361. Which law states entropy of all pure crystalline solids is zero at absolute zero ?

- A. First law of thermodynamics
- B. Second law of thermodynamics
- C. Third law of thermodynamics
- D. Hess's law

Answer: B

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362. The enthalpies of formation of $C_2H_2(g)$ and $C_6H_6(g)$ at 298 K are 230 and 85 kJ/mol respectively. The enthalpy change for the reaction $3C_2H_2(g) \rightarrow C_6H_6(g)$ is

- A. 605 kJ/mol
- B. 865 kJ/mol
- C. -605 kJ/mol
- D. -865 kJ/mol

Answer: C

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363. For an ideal gas, the Joule Thomson coefficient is

- A. Zero
- B. negative
- C. positive
- D. depend on molecular weight

Answer: C

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364. The relation of ΔH and ΔE is represented as

A. $\Delta H = \Delta E - P\Delta V$

B. $\Delta E = \Delta H - P\Delta V$

C. $\Delta H = \Delta E + \Delta n_g RT$

D. $\Delta E = \Delta V + \Delta H$

Answer: A

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365. Calculate the temperature at which $\Delta G = -5.2 \text{ kJ mol}^{-1}$, $\Delta H = 145.6 \text{ kJ mol}^{-1}$ and $\Delta S = 216 \text{ JK}^{-1} \text{ mol}^{-1}$ for a chemical reaction

A. 698.1°C

B. 698.1K

C. 130K

D. 130°C

Answer: C



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366. When ice melts into water entropy

- A. becomes zero
- B. decreases
- C. increase
- D. remains the same

Answer: C



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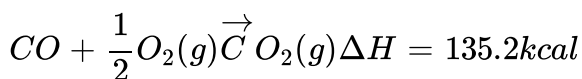
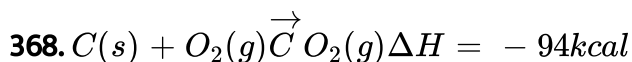
367. Which of the following is not the extensive property?

- A. Gibbs free energy
- B. Enthalpy
- C. Entropy

D. Viscosity

Answer: C

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The heat of formation of CO(g) is ?

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369. In order to decompose 9g water 142.5 kJ heat is required. Hence, the enthalpy of formation of water is

A. -142.5 kJ

B. +142.5kJ

C. -285 kJ

D. $+285 \text{ kJ}$

Answer: A

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370. The enthalpy of combustion of $C_6H_6(l)$ is -3250 kJ . When 0.39 g of benzene is burnt in excess of oxygen in an open vessel, the amount of heat liberated is

A. 16.26 J

B. -16.25 J

C. 32.5 J

D. 32.5 kJ

Answer: C

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371. For the reaction $H_2O(s) \rightleftharpoons H_2O(l)$ at $0^\circ C$ and normal pressure

A. $\Delta H > T\Delta S$

B. $\Delta H = T\Delta S$

C. $\Delta H = \Delta G$

D. $\Delta H < T\Delta S$

Answer: B



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372. $H_2 + Cl_2 \xrightarrow{2} HCl$, $\Delta H = 194kJ$ In this reaction heat of formation of HCl in kJ is

A. $+914kJ$

B. $+97kJ$

C. $-97kJ$

D. $-194 kJ$

Answer: B

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373. $N_2(g) + 3H_2(g) \rightarrow 2NH_3(g)$ What is the enthalpy of the above reaction when the enthalpy formation of ammonia is $-46kJmol^{-1}$?

- A. 184 kJ
- B. -184 kJ
- C. 46 kJ
- D. -46 kJ

Answer: B

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374. $N_2(g) + 3H_2(g) \rightarrow 2NH_3(g)$. For the above reaction the correct statement is

A. $\Delta H < \Delta E$

B. $\Delta H = \Delta E$

C. $\Delta H > \Delta E$

D. None of these

Answer: A

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375. A chemical process is carried out in a thermostat maintained at $25^{\circ}C$, it is known as

A. isothermal

B. isobaric

C. adiabatic

D. isoentropic

Answer: A

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376. Identify the correct statement from the following in a chemical reaction.

- A. The entropy always increases
- B. The change in entropy along with suitable change in enthalpy decides the rate of reaction
- C. The enthalpy always decreases
- D. Both the enthalpy and the entropy remain constant

Answer: B

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377. The incorrect expression among the following is

A.
$$\frac{\Delta G_{system}}{(\Delta S_{\rightarrow tal})} = T$$

B. In isothermal process, $W_{reversib} = -nRt \ln \frac{V_f}{V_i}$

C. $\ln K = \frac{\Delta H^\circ - T\Delta S^\circ}{RT}$

D. $K = e^{\frac{-\Delta G}{RT}}$

Answer: C

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378. A sample of liquid in a thermally insulated container is stirred for 1h by a mechanical attachment to a motor in the surroundings, which of the following thermodynamic quantity for the system is zero ?

- A. Work (W)
- B. Change in internal energy (ΔE)
- C. Change in enthalpy (ΔH)
- D. None of the above

Answer: D



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379. For an isolated system, $\Delta U = 0$, then

- A. $\Delta S = 0$
- B. $\Delta S < 0$
- C. $\Delta S > 0$
- D. The value of ΔS cannot be predicted

Answer: A



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380. A gas undergoes a thermodynamical process but the volume of the gas does not change. What is the nature of this process?

- A. Isothermal
- B. Adiabatic

C. Isobaric

D. Isochoric

Answer: D

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381. Among the following for spontaneity of chemical reaction there should be

A. decrease in entropy and increase in free energy

B. decrease in entropy and free energy both

C. increase in entropy and decrease in free energy

D. increase in entropy and free energy both

Answer: C

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382. Which of the following processes is associated with decrease in entropy ?

- A. Vaporisation of a mole of water into steam at its b.p
- B. Dissociation of a mole of common salt in water at 300K
- C. Mixing of two partially miscible liquids
- D. Crystallisation of a salt from its saturated solution

Answer: D



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383. Which of the following statements is true?

- A. The total entropy of the universe is continuously decreasing.
- B. The total energy of the universe is continuously decreasing.
- C. The total energy of the universe remains constant.
- D. The total entropy of the universe remains constant.

Answer: C

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384. A coffee cup calorimeter initially contains 125g of water at a temperature of 24.2°C . After adding 10.59 of KBr, the temperature becomes 21.1°C . The heat of solution is

A. 40 J/g

B. 167 J/g

C. 155 J/g

D. 420.05 J/g

Answer: B

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385. One mole of ice is converted into water at 273K. The entropies of $H_2O(s)$ and $H_2O(l)$ are 38.20 and $60.01 \text{ J mol}^{-1} \text{ K}^{-1}$ respectively. The enthalpy change for the conversion is .

A. 3 kJ mol^{-1}

B. 4 kJ mol^{-1}

C. 5 kJ mol^{-1}

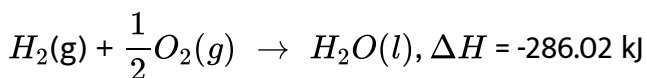
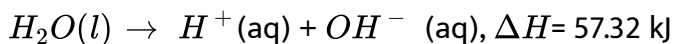
D. 6 kJ mol^{-1}

Answer: D



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386. Given that



Then calculate the enthalpy of formation of OH^- at 25°C .

A. -228.8 kJ

B. -343.53 kJ

C. +228.8 kJ

D. +343.52 kJ

Answer: A

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387. Calculate the amount of heat evolved when 500cm^3 of 0.1 M HCl is mixed with 200cm^3 of 0.2 M NaOH.

A. 57.3 kJ

B. 2.865 kJ

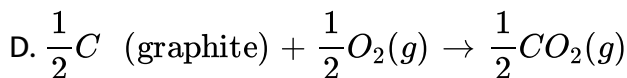
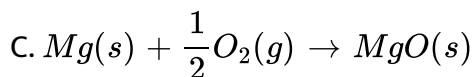
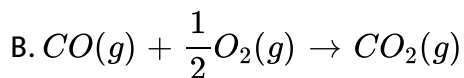
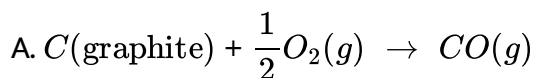
C. 2.292 kJ

D. 0.573 kJ

Answer: C

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388. In which of the following reactions, standard reaction entropy changes (ΔS°) is positive and standard Gibbs energy change (ΔG°) decreases sharply with increasing temperature?



Answer: A

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389. The enthalpy of fusion of ice is 1.435 kcal/mol. The molar entropy change for the melting of ice at $0^\circ C$ is

A. 10.52 cal/(mol K)

B. 21.04 cal/(mol K)

C. 5.260 cal/(mol K)

D. 0.526 cal/(mol K)

Answer: C



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390. Standard enthalpy of vaporisation, $\Delta_{vap}H^\theta$ for water at $100^\circ C$ is $40.66 kJ mol^{-1}$. The internal energy of vaporisation of water at $100^\circ C$ (in $kJ mol^{-1}$) is (Assume water vapour to behave like an ideal gas).

A. + 37.56

B. -43.76

C. + 43.76

D. + 40.66

Answer: A



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391. Equal volumes of two monoatomic gases, A and B, at same temperature and pressure are mixed. The ratio of specific heats $\left(\frac{C_p}{C_v}\right)$ of the mixture will be

A. 0.83

B. 1.5

C. 3.3

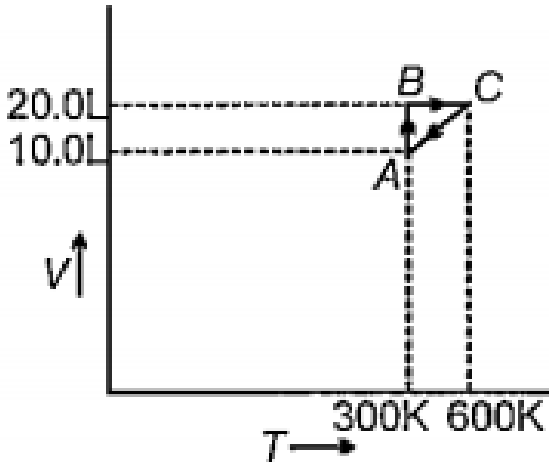
D. 1.67

Answer: D



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392. This graph expresses the various steps of the system containing 1 mol of gas. Which type of process system has when it moves from C to A?



- A. Isochoric
- B. Isobaric
- C. Isothermal
- D. Cyclic

Answer: B



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393. Which of the following is correct ?

- A. Evaporation of water causes an increase in disorder of the system
- B. Melting of ice causes a decrease in randomness of the system
- C. Condensation of steam causes an increase in disorder to the system
- D. There is practically no change in the randomness of the system when water is evaporated

Answer: A

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394. On passing C ampere of current for time ' t ' sec through 1L of 2 (M) $CuSO_4$ solution (atomic weight of Cu 63.5), the amount ' m ' of Cu (in g) deposited on cathode will be

A. $m = C \frac{t}{63.5 \times 96500}$

$$B. m = C \frac{t}{31.75 \times 96500}$$

$$C. m = \frac{C \times 96500}{31.75 \times t}$$

$$D. m = 31.75 \times C \times t / (96500)$$

Answer: D



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395. Entropy of the universe is

A. constant

B. zero

C. continuously decreasing

D. continuously increasing

Answer: D



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396. The equilibrium constant of a reaction is 0.008 at 298 K. The standard free energy change of the reaction at the same temperature is

- A. -11.96 kJ
- B. -5.43 kJ
- C. -8.46 kJ
- D. +11.96 kJ

Answer: D



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397. A gas undergoes a thermodynamical process but the volume of the gas does not change. What is the nature of this process?

- A. Isothermal
- B. Adiabatic
- C. Isobaric

D. Isochoric

Answer: D

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398. The occurrence of reaction is impossible if

A. ΔH is $+ve$, ΔS is also $+ve$

B. ΔH is $-ve$, ΔS is also $-ve$

C. ΔH is $-ve$, ΔS is $+ve$

D. ΔH is $+ve$, ΔS is also $-ve$

Answer: D

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399. The bond dissociation energies of XY , X_2 and Y_2 (all diatomic molecules) are in the reaction 1:1:0:5 and ΔH_r for the formation of XY is -200 kJ mol^{-1} what will be the bond dissociation energy of X_2 ?

A. 400 kJ mol^{-1}

B. 300 kJ mol^{-1}

C. 200 kJ mol^{-1}

D. 800 kJ mol^{-1}

Answer: D



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400. Work done during isothermal expansion of one mole of an ideal gas from 10 atm to 1 atm at 300 K is

A. 4938.8 J

B. 4138.8 J

C. 5744.1 J

D. 6257.2 J

Answer: C

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401. The amount of the heat released when 20 ml 0.5M NaOH is mixed with 100 mL 0.1 M HCl is x kJ. The heat of neutralisation is

A. $-100 \times \text{kJ/mol}$

B. $-50 \times \text{kJ/mol}$

C. $+100x \text{ k} \frac{\text{J}}{\text{m}} \text{ol}$

D. $+50x \text{ kJ/mol}$

Answer: A

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402. A plot of $\ln k$ against $1/T$ (abscissa) is expected to be a straight line with intercept on coordinate axis equal to

A. $\frac{\Delta S^\circ}{2.303R}$

B. $\Delta \frac{S}{R}$

C. $-\frac{\Delta S}{R}$

D. $R \times \Delta S^\circ$

Answer: B



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403. Based on the first law of thermodynamics, which one of the following is correct

A. For an isochoric process $\Delta E = -Q$

B. For an adiabatic process $\Delta E = -W$

C. For an isothermal process $Q = -W$

D.

Answer: D

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404. The value of enthalpy change (ΔH) for the reaction $C_2H_5OH(l) + 3O_2(g) \rightarrow 2CO_2(g) + 3H_2O(l)$, at $27^\circ C$ is -1366.5 kJ .

The value of internal energy change for the above reaction at this temperature will be

A. -1371.5 kJ

B. -1369.0 kJ

C. -1364.0 kJ

D. -1361.5 kJ

Answer: C

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405. Consider the reaction, $4NO_2(g) + O_2(g) \rightleftharpoons 2N_2O_5(g)$, $\Delta_r H = -111kJ$. If $N_2O_5(s)$ is formed instead of $N_2O_5(g)$ in the above reaction, the $\Delta_r H$ value will be (Given, ΔH of sublimation for N_2O_5 is $54kJmol^{-1}$)

- A. -165 kJ
- B. +54kJ
- C. +219kJ
- D. -219 kJ

Answer: D

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406. Molar heat capacity of aluminum is $25JK^{-1}mol^{-1}$. The heat necessary to raise the temperature of 54g of aluminium (Atomic mass $27g mol^{-1}$) from $30^\circ C$ to $50^\circ C$ is

(1) 1.5 kJ

(2) 0.5 kJ

(3) 1.0 kJ

(4) 2.5 kJ

(5) 2.0 kJ



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407. The entropy change involved in the isothermal reversible expansion of 2 moles of an ideal gas from a volume of 10dm^3 to a volume of 100dm^3 at 27°C is

A. $38.3\text{Jmol}^{-1}\text{K}^{-1}$

B. $38.8\text{Jmol}^{-1}\text{K}^{-1}$

C. $32.3\text{Jmol}^{-1}\text{K}^{-1}$

D. $42.3\text{Jmol}^{-1}\text{K}^{-1}$

Answer: A



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408. In view of the signs of $\Delta_r G^\circ$ for the following reactions $\text{PbO}_2 + \text{Pb} \rightarrow 2\text{PbO}$, $\Delta_r G^\circ < 0$, $\text{SnO}_2 + \text{Sn} \rightarrow 2\text{SnO}$, $\Delta_r G^\circ > 0$ Which oxidation states are more characteristic for lead and tin?

- A. For lead +4, for tin +2
- B. For lead +2, for tin +2
- C. For lead +4, for tin +4
- D. For lead +2, for tin +4

Answer: D

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409. For the reversible reaction, $A(s) + B(g) \rightleftharpoons C(g) + D(g)$, $\Delta_r G^\circ = -350\text{kJ}$, which one of the following statements is true?

- A. The reaction is thermodynamically non feasible.

- B. The entropy change is negative.
- C. Equilibrium constant is greater than one.
- D. The reaction should be instantaneous.

Answer: C

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410. Carbon cannot reduce Fe_2O_3 to Fe at a temperature below 983 K because

- A. free energy change for the formation of CO is more negative than that of Fe_2O_3 ,
- B. CO is thermodynamically more stable than Fe_2O_3 ,
- C. carbon has higher affinity towards oxygen than iron
- D. iron has higher affinity towards oxygen than carbon

Answer: C

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411. The enthalpy of vaporisation of benzene is $+35.3 \text{ kJ/mol}$ at its boiling point of 80°C . The entropy change in the transition of vapour to liquid at its boiling point is (in $\text{J mol}^{-1} \text{K}^{-1}$).

- A. -100
- B. +100
- C. +342
- D. -342

Answer: A

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412. A container of 1.0 L capacity filled with 1.0 mole of ideal gas is connected to an evacuated vessel of 9.0L. Calculate change in entropy. ($R = 1.987 \text{ cal.}$)

A. $0.188 \text{ cal. } K^{-1} \text{ mol}^{-1}$

B. $0.4576 \text{ cal. } K^{-1} \text{ mol}^{-1}$

C. $4.576 \text{ cal. } K^{-1} \text{ mol}^{-1}$

D. $4.366 \text{ cal. } K^{-1} \text{ mol}^{-1}$

Answer: D

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413. Which of the following is the correct option for free expansion of an ideal gas under adiabatic condition ?

A. $q = 0, \Delta T < 0, w \neq 0$

B. $q = 0, \Delta T \neq 0, w = 0$

C. $q \neq 0, \Delta T = 0, w = 0$

D. $q = 0, \Delta T = 0, w = 0$

Answer: D

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414. Enthalpy change for the reaction, $4H(g) \rightarrow 2H_2(g)$ is -869.5 kJ. The dissociation energy of H-H bond is

A. $+217.4$ kJ

B. -434.8 kJ

C. -868.6 kJ

D. $+434.8$ kJ

Answer: D

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415. If the enthalpy change for the transition of liquid water to steam is 300 kJ mol⁻¹ at $27^\circ C$, the entropy change for the process would be

A. 1000 JK⁻¹ mol⁻¹

B. $10JK^{-1}mol^{-1}$

C. $1.0JK^{-1}mol^{-1}$

D. $0.1JK^{-1}mol^{-1}$

Answer: A

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416. Based on the first law of thermodynamics, which one of the following is correct

A. For an isothermal process, $q = +w$

B. For an isochoric process, $\Delta U = -q$

C. For an adiabatic process, $\Delta U = -w$

D. For a cyclic process, $q = -w$

Answer: D

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417. The enthalpy of solution of sodium chloride is 4kJmol^{-1} and its enthalpy of hydration of ions is -784kJmol^{-1} . Then the lattice enthalpy of NaCl (in kJ mol^{-1}) is

(1) + 788

(2) + 4

(3) + 398

(4) + 780

+ 394



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418. Four grams of graphite is burnt in a bomb calorimeter of heat capacity 60kJK^{-1} in excess of oxygen at 1 atm pressure. The temperature rises from 300 to 304 K, What is the enthalpy of combustion of graphite (in kJ mol^{-1})

(1) 360

(2) 1440

(3) -720

(4) -1440

(5) -520



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419. Which reaction, with the following values of ΔH , ΔS at 400K is spontaneous and endothermic ?

A. $\Delta H = -48\text{kJ}$, $\Delta S = +135\frac{\text{J}}{\text{K}}$

B. $\Delta H = -48\text{kJ}$, $\Delta S = -135\frac{\text{J}}{\text{K}}$

C. $\Delta H = +48\text{kJ}$, $\Delta S = +135\frac{\text{J}}{\text{K}}$

D. $\Delta H = +48\text{kJ}$, $\Delta S = -135\frac{\text{J}}{\text{K}}$

Answer: C



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420. The enthalpy of vaporisation of benzene is $+35.3 \text{ kJ/mol}$ at its boiling point of 80°C . The entropy change in the transition of vapour to liquid at its boiling point is (in $\text{J mol}^{-1} \text{K}^{-1}$).

- A. -100
- B. +100
- C. +342
- D. -342

Answer: A



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421. For the reversible reaction,
 $A(s) + B(g) \rightleftharpoons C(g) + D(g)$, $\Delta G^\circ = -350 \text{ kJ}$, which one of the following statement is true ?

- A. The reaction is thermodynamically non feasible.

- B. The entropy change is negative.
- C. Equilibrium constant is greater than one.
- D. The The reaction should be instantaneous.

Answer: C

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422. A reaction is spontaneous at low temperature but non spontaneous at high temperature. Which of the following

- (1) $\Delta H > 0, \Delta S > 0$
- (2) $\Delta H < 0, \Delta S < 0$
- (3) $\Delta H > 0, \Delta S = 0$
- (4) $\Delta H < 0, \Delta S < 0$
- (5) $\Delta H = 0, \Delta S < 0$

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423. At the sublimation temperature, for the process $CO_2(s) \rightleftharpoons CO_2(g)$

(1) ΔH , ΔS and ΔG are all positive (2) $\Delta H > 0$, $\Delta S > 0$ and $\Delta G < 0$

(3) $\Delta H < 0$, $\Delta S > 0$ and $\Delta G < 0$

(4) $\Delta H > 0$, $\Delta S > 0$ and $\Delta G > 0$

(5) $\Delta H > 0$, $\Delta S > 0$ and $\Delta G = 0$



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424. Match the thermodynamic processes given under Column I with the expression given under Column II.

Column I	Column II
(A) Freezing of water at 273 K and 1 atm	(P) $q = 0$
(B) Expansion of 1 mol of an ideal gas into a vacuum under isolated conditions	(Q) $w = 0$
(C) Mixing of equal volumes of two ideal gases at constant temperature and pressure in an isolated container	(R) $\Delta S_{\text{sys}} = 0$
(D) Reversible heating of $\text{H}_2(\text{g})$ at 1 atm from 300 K to 600 K, followed by reversible cooling to 300 K at 1 atm	(S) $\Delta U = 0$
	(T) $\Delta G = 0$



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425. For the process $\text{H}_2\text{O}(l) \rightarrow \text{H}_2\text{O}(g)$ at $T = 100^\circ\text{C}$ and 1 atmosphere pressure, the correct choice is

A. $\Delta S_{system} > 0$ and ΔS (surroundings) > 0

B. $\Delta S_{system} > 0$ and ΔS (surroundings) < 0

C. $\Delta S_{system} < 0$ and ΔS (surroundings) > 0

D. $\Delta S_{system} < 0$ and ΔS (surroundings) < 0

Answer: B

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426. For complete combustion of ethanol,

$C_2H_5OH(l)$, + $3O_2(g) \rightarrow 2CO_2(g) + 3H_2O(l)$, the amount of heat produced as measured in bomb calorimeter, is $1364.47kJmol^{-1}$ at $25^\circ C$.

Assuming ideality the Enthalpy of combustion, $\Delta_c H$, for the reaction will be:

($R = 8.314Jmol^{-1}$)

A. $-1366.95kJmol^{-1}$

B. $-1361.95kJmol^{-1}$

C. $-1460.95 \text{ kJ mol}^{-1}$

D. $-1350.95 \text{ kJ mol}^{-1}$

Answer: A

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427. The standard enthalpies of formation of $\text{CO}_2(g)$, $\text{H}_2\text{O}(l)$ and glucose (s) at 25°C are -400 kJ/mol , -300 kJ/mol and -1300 kJ/mol , respectively. The standard enthalpy of combustion per gram of glucose at 25°C is

A. $+2900 \text{ kJ}$

B. -2900 kJ

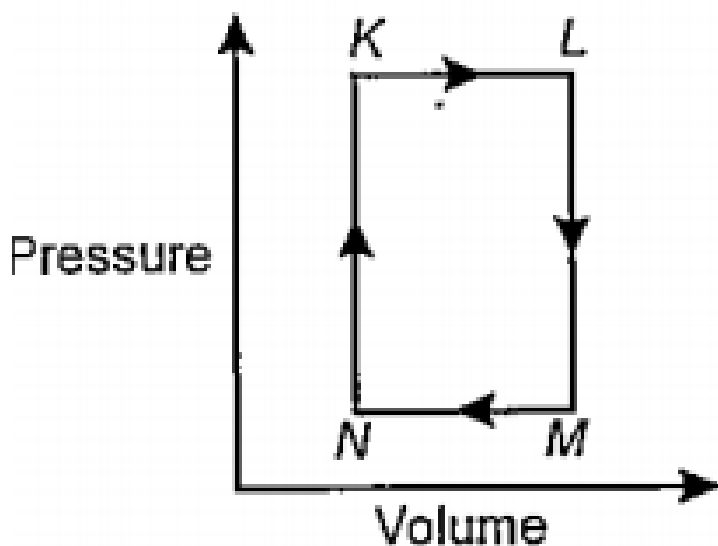
C. -16.11 kJ

D. $+16.11 \text{ kJ}$

Answer: C

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428. A fixed mass m of a gas is subjected to transformation of states from K to L to M to N and back to K as shown in the figure.



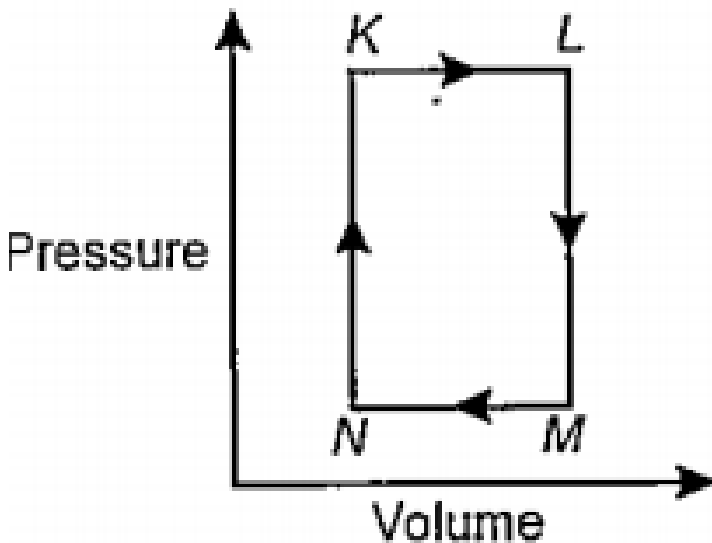
The pair of Isochoric processes among the transformation of states is

- A. heating, cooling, heating, cooling
- B. cooling, heating, cooling, heating
- C. heating, cooling, cooling, heating
- D. cooling, heating, heating, cooling

Answer: C

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429. A fixed mass m of a gas is subjected to transformation of states from K to L to M to N and back to K as shown in the figure.



The pair of Isochoric processes among the transformation of states is

- A. K to L and L to M
- B. L to M and N to K
- C. L to M and M to N

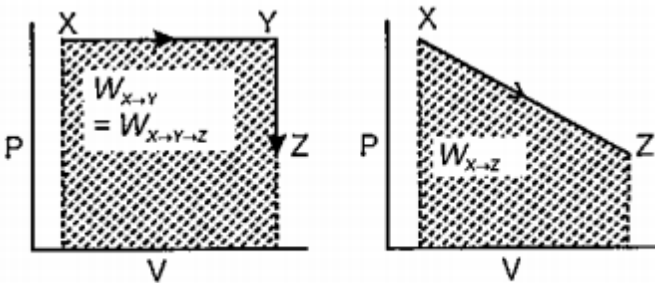
D. M to N and N to K

Answer: B

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430. For an ideal gas, consider only p-V work in going from an initial state X to the final state Z. The final state Z can be reached by either of the two paths shown in the figure. Which of the following choice(s) is (are) correct

?



[Take ΔS as change in entropy and W as work done].

A. $\Delta S_{X \rightarrow Z} = \Delta S_{X \rightarrow Y} + \Delta S_{Y \rightarrow Z}$

B. $W_{X \rightarrow Z} = W_{X \rightarrow Y} + W_{Y \rightarrow Z}$

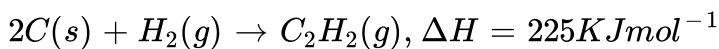
C. $W_{X \rightarrow Y \rightarrow Z} = W_{X \rightarrow Y}$

$$D. \Delta S_{X \rightarrow Y \rightarrow Z} = \Delta S_{X \rightarrow Y}$$

Answer: A::C

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431. Using the data provided calculate the multiple bond energy ($KJmol^{-1}$) of a C=C bond in C_2H_2 consider the bond energy of a C-H bond is $350KJmol^{-1}$



A. 1165

B. 837

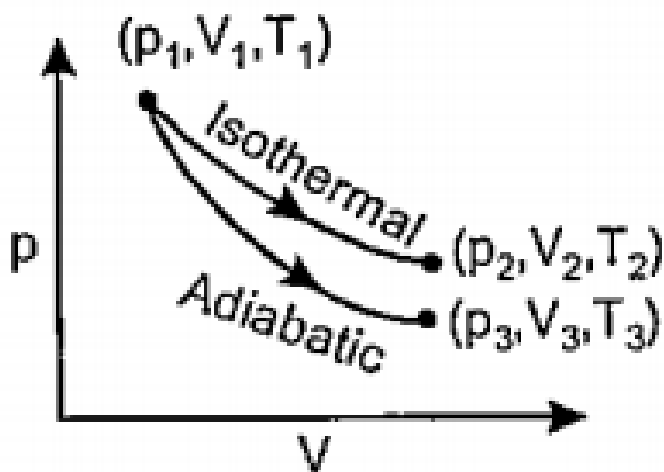
C. 865

D. 815

Answer: D



432. The reversible expansion of an ideal gas under adiabatic and isothermal conditions is shown in the figure. Which of the following statement(s) is (are) correct ?



A. $T_1 = T_2$

B. $T_3 > T_1$

C. $W_{isothermal} > W_{adiabatic}$

D. $\Delta U_{isothermal} > \Delta U_{adiabatic}$

Answer: A::C::D

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433. A gas expands from a volume of $1m^3$ to a volume of $2m^3$ against an external pressure of $10^5 Nm^{-2}$. the work done by the gas will be

A. $10^4 J$

B. $10^2 J$

C. $10^3 J$

D. $10^5 J$

Answer: D

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