

# **CHEMISTRY**

# **BOOKS - NTA MOCK TESTS**

# **THERMODYNAMICS TEST-1**

# **Single Choice**

1. The S - S bond energy, if

$$\Delta H_f^{\,\circ}(E_t-S-E_t)=\,-\,147$$
 mol,

ol, and

$$\Delta H_f^{\,\circ}(E_t-S-S-E_t)=\,-\,202$$
 kJ/mol

$$\Delta H_f^{\,\circ} S(g) = \, + \, 233$$
 kJ/mol, is:

A. 268 kJ

- B. 126 kJ
- C. 278 kJ
- D. 572 kJ

#### **Answer: C**



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**2.** The lattice energy of solid NaCl is 180 kcal/mol. The dissolution of the solid in water in the form of ions is endothermic to the extent of 1 kcal/mol. If the hydration energies of  $Na^+$  and  $Cl^-$  are in the ratio 6:5, what is the enthalpy of hydration of Nation ?

A. -8.5 kcal  $mol^{-1}$ 

B.  $-97.64 \text{ kcal mol}^{-1}$ 

 $\mathsf{C.} + 82.6 \, \mathsf{kcal} \, \mathrm{mol}^{-1}$ 

D.  $+100 \text{ kcal mol}^{-1}$ 

# **Answer: B**



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**3.** Diborane is a potential rocket fuel which undergoes combustion according to the reaction:

$$B_2H_6(g) + 3O_2(g) o B_2O_3(s) + 3H_2O(g)$$

From the following data, calculate the enthalpy change

for the combustion of diborane.

A. 
$$-2035kJmol^{-1}$$

B. 
$$-2045 \text{kj mol}^{-1}$$

$$C.-2135 kJ \text{ mol}^{-1}$$

$$D.-2285 \mathrm{kJ} \; \mathrm{mol}^{-1}$$

#### **Answer: A**



**4.** 2 mol of an ideal gas at 27°C temperature is expanded reversibly from 2 L to 20 L. Find the entropy change. (R = 2 cal / mol K)

A. 
$$9.2calK^{-1}$$

B. 
$$92.1calK^{-1}$$

C. 
$$46calK^{-1}$$

D. 
$$4.6calK^{-1}$$

# **Answer: A**



**5.** The enthalpy of neutralisation of  $NH_4OH$  and  $CH_3COOH$  is -10.5 kcal  $\mathrm{mol}^{-1}$  and enthalpy of neutralisation of  $CH_3COOH$  with strong base is -12.5 kcal  $\mathrm{mol}^{-1}$ . The enthalpy of ionisation of  $NH_4OH$  will be:

- A. 4.0kcal mol $^{-1}$
- B. 3.0kcal mol<sup>-1</sup>
- C. 2.0kcal mol $^{-1}$
- D. 3.2kcal mol<sup>-1</sup>

# **Answer: C**



- 6. The enthalpy of hydrogenation of cyclohexene is
- $-119.5 \mathrm{kJ} \; \mathrm{mol}^{-1}.$  If resonance energy of benzene is
- $-150.4 \mathrm{kJ} \; \mathrm{mol}^{-1}$  its enthalpy of hydrogenation would be
  - $A. -358.5 kJ \text{ mol}^{-1}$
  - $B. -508.9 kJ \text{ mol}^{-1}$
  - $C. -208.1 kJ \text{ mol}^{-1}$
  - $D. 269.9 kJ \text{ mol}^{-1}$

#### **Answer: C**



**7.** The bond dissociation energies of gaseous  $H_2,\,Cl_2$  and HCl are 104, 58 and 103 kcal, respectively. The enthalpy of formation of HCl gas would be:

- $\mathsf{A.}-44\ \mathsf{kcal}$
- $\mathrm{B.}-88~\mathrm{kcal}$
- $\mathsf{C.}-22~\mathsf{kcal}$
- D.-11 kcal

# **Answer: C**



**8.** The change in entropy of 2 moles of an ideal gas upon isothermal expansion at 243.6 K from 20 L to the state where pressure becomes 1 atm is (Given: In 2 = 0.693)

- A. 1.385 cal/K
- $\mathrm{B.}-1.2~\mathrm{cal/K}$
- C. 1.2 cal/K
- D. 2.77 cal/K

## **Answer: D**



**9.** The work done in an open vessel at 300 K, when 112 g iron reacts with dil. HCl is:

$$\Big(R=2 ext{cal mol}^{\,-1}K^{\,-1}\Big)$$

- A. 1200 cal
- B. 600 cal
- C. 300 cal
- D. 200 cal

**Answer: A** 



# 10. At 1000 K, from the data,

$$N_2(g) + 3H_2(g) 
ightarrow 2NH_3(g), \Delta_r H^{\,\circ} = \, -\, 123.77 {
m kJ \; mol}^{-\,1}$$

Substance	N <sub>2</sub>	$H_2$	NH <sub>3</sub>	
C <sub>p</sub> /R	3. 5	3. 5	4	

The heat of formation of ammonia in  ${
m kJ\ mol}^{-1}$  at 300 K

$$A. - 88.85$$

is:

$$B. + 88.85$$

$$C. -44.42$$

$$D. + 44.42$$

#### **Answer: C**



11. Enthalpy of atomization of  $C_2H_6(g)$  and  $C_3H_8(g)$  are 620 and  $880kJ\mathrm{mol}^{-1}$  respectively. The C-C and C-H. bond energies are respectively.

- A. 80 and 60  ${
  m kJ~mol}^{-1}$
- B. 80 and 90  ${
  m kJ~mol}^{-1}$
- C. 70 and 90  ${
  m kJ~mol}^{-1}$
- D. 200 and 80  ${
  m kJ~mol}^{-1}$

## **Answer: B**



12. On the basis of the following thermochemical data:

$$igl[ \Delta_f H^{\,\circ} \;\; ext{ of } \;\; H^{\,+}(aq) = 0 igr]$$

$$H_2O(l)
ightarrow H^+(aq)+OH^-(aq), \Delta H^\circ = 57.32kJ$$

$$H_2(g) + rac{1}{2} O_2(g) o H_2 O(l), \Delta H^{\,\circ} = \, -\, 286.02 \, ext{kJ}$$

What is the value of enthalpy of formation of  $OH^-$  at 25°C?

$$\mathsf{A.}-22.88~\mathsf{kJ}$$

$$\mathrm{B.}-228.88~\mathrm{kJ}$$

$$\mathrm{C.} + 228.88~\mathrm{kJ}$$

$$\mathrm{D.}-343.52~\mathrm{kJ}$$

## **Answer: B**



**13.** The free energy change for the following reactions are

given below:

$$C_2 H_2(g) + rac{5}{2} O_2(g) 
ightarrow 2 C O_2(g) + H_2 O(l)$$
 ,

$$\Delta G^{\circ} = \, -\, 1234 \, \mathrm{kJ}$$

$$C(s) + O_2(g) 
ightarrow CO_2(g)$$
 ,

$$\Delta G^{\,\circ} = \,-\,394\,\mathrm{kJ}$$

$$H_2(g)+rac{1}{2}O_2(g)
ightarrow H_2O(l)$$
 ,

$$\Delta G^{\circ} = \, -\, 237 \, ext{kJ}$$

What is the standard free energy change for the following reaction?

$$H_2(g)+2C(s)
ightarrow C_2H_2(g)$$
 ?

$$\mathrm{A.}-209~\mathrm{kJ}$$

$$\mathrm{B.}-2259~\mathrm{kJ}$$

$$\mathrm{C.} + 2259~\mathrm{kJ}$$

D. 209 kJ

## **Answer: D**



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**14.** 1 mole of  $H_2SO_4$  is mixed with 2 moles of NaOH. The heat evolved will be:

A. 57.3 kJ

B. 2 imes 57.3 kJ

 $\mathsf{C.}\,57.3/2\,\mathsf{kJ}$ 

D. Cannot be predicted

# **Answer: B**



**15.** The enthalpy of combustion of  $H_2$ , cyclohexene ( $C_6H_{10}$ ) and cyclohexane ( $C_6H_{12}$ ) are -241, -3800 and  $-3920kJ\mathrm{mol}^{-1}$  respectively. The heat of hydrogenation of cyclohexene is:

- A. 121kJ mol $^{-1}$
- B.  $-121 \text{kJ mol}^{-1}$
- $C. + 242 kJ \text{ mol}^{-1}$
- D.  $-242 \text{kJ mol}^{-1}$

# **Answer: B**



**16.** The heat of atomisation of  $PH_3$  (g) is 228 kcal  $\mathrm{mol}^{-1}$  and that of  $P_2H_4$  (g) is 335 kcal  $\mathrm{mol}^{-1}$ . The energy of P-P bond is:

- A. 102 kcal  $\mathrm{mol}^{-1}$
- B. 31 kcal  $\mathrm{mol}^{-1}$
- C. 26 kcal  $\mathrm{mol}^{-1}$
- D. 204 kcal  $\mathrm{mol}^{-1}$

# **Answer: B**



17. At what minimum pressure (in terms of kPa) of given volume of an ideal gas ( $C_{p.m}=\frac{7}{2}R$ ), originally at 400 K and 100 kPa pressure, be irreversibly adiabatically compressed in order to raise its temperature to 600 K?

- A. 362.5 kPa
- B. 275.0 kPa
- C. 437.5 kPa
- D. 550.0 kPa

#### **Answer: B**



**18.** 16 g oxygen gas expands at STP to occupy double of its original volume. The magnitude of work done during the process is:

- A. 260 cal
- B. 180 cal
- C. 130 cal
- D. 271.6 cal

# **Answer: D**



**19.** 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, AH $^{\rm o}$  of neutralization of a strong acid and a strong base is 140 kcal  ${
m mol}^{-1}$ .

- A. 14 kcal
- B. 35 cal
- C. 10 cal
- D. 7.5 cal

## **Answer: B**



**20.** If  $\Delta_f H^\circ(C_2H_4)$  and  $\Delta_f H^\circ(C_2H_6)$  are a and b  $m kcal\ mol^{-1}$ , then, the heat of hydrogenation of  $C_2H_4$  IS:

A. 
$$a+b$$

B. 
$$a-b$$

$$C.b-a$$

D. 
$$a-2b$$

#### **Answer: C**



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21. The enthalpies of neutralization BOH and a strong base BOH by HCl are – 12250 cal / mol and - 13000 cal /

mol respectively. When one mole of HCl is added to a solution containing 1 mole of AOH and 1 mole of BOH, the enthalpy change was - 12500 cal / mol. In what ratio is the acid distributed between AOH and BOH respectively.

- A.2:1
- B. 2:3
- C. 1: 2
- D. 3:2

#### **Answer: A**



**22.** In the conversion of limestone to lime,  $CaCO_3(s)+CaO(s)+CO_2$  (g), the values of  $\Delta H^\circ$  and  $\Delta S^\circ$  are  $179.1 \mathrm{kJ} \ \mathrm{mol}^{-1}$  and  $160.2 J K^{-1} \mathrm{mol}^{-1}$  respectively at 298 K and 1 bar. Assuming  $\Delta H^\circ$  and  $\Delta S^\circ$  remain constant with temperature, at which minimum temperature does the conversion of limestone to lime will be spontaneous ?

A. 1118 K

B. 1008 K

C. 1200 K

D. 845 K

#### **Answer: A**

**23.** For an ideal gas,  $\frac{C_{
m p.m}}{C_{
m v.m}}=\gamma$ . The molecular mass of a gas is M, its specific heat capacity at constant volume is:

A. 
$$rac{R}{M(\gamma-1)}$$

B. 
$$\frac{M}{R(\gamma-1)}$$

C. 
$$rac{\gamma RM}{\gamma-1}$$

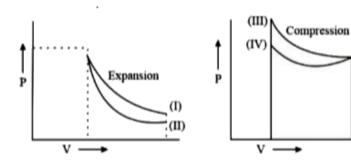
D. 
$$\frac{\gamma R}{M(\gamma-1)}$$

## **Answer: A**



# 24. Which of the figures given below show the adiabatic

process?



A. I,III

B. I,III

C. II, IV

D. I,IV

# Answer: A



**25.** The value of  $\log_{10}$  K for a reaction A o B is Given:

$$\Delta_r H_{298K}^{\,\circ} = \,-\,54.07 k J \mathrm{mol}^{\,-\,1}$$
 ,

$$\Delta_r S_{298K}^{\,\circ} = 10 J K^{\,-1} \mathrm{mol}^{\,-1}$$

$$R = 8.314 J K^{-1} \text{mol}^{-1}$$

$$2.303 \times 8.314 \times 298 = 5705$$

A. 5

B. 10

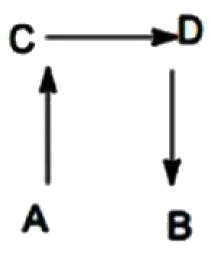
C. 95

D. 100

#### **Answer: B**



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



Given:

$$\Delta S_{A 
ightarrow C} = 50$$
 c.u,

$$\Delta S_{C o D} = 30$$
 c.u.

 $\Delta S_{B 
ightarrow D} = 20$  c.u where e.u is the entropy unit.

Then,  $\Delta S_{A 
ightarrow B}$  is:

A.+60 eu

- B. + 100 e.u
- $\mathsf{C.}-60\,\mathsf{e.u}$
- D. 100 e.u

## **Answer: A**



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**27.** If  $\Delta H$  is the change in enthalpy and  $\Delta E$ , the change in internal energy accompanying a gaseous reaction, then:

A.  $\Delta H$  is always greater than  $\Delta E$ 

B.  $\Delta H < \Delta E$  only if the number of moles of the products is greater than the number of moles of the reactants.

- C.  $\Delta H$  is always less than  $\Delta E$
- D.  $\Delta H < \Delta E$  only if the number of moles of products is less than the number of moles of the reactants.

#### **Answer: D**



**28.** One mole of an ideal gas at 300 K is expanded isothermally from an initial volume of 1 litre to 10 litre.

 $\Delta E$  for this process is:  $(\mathsf{R} = \mathsf{2} \; \mathsf{cal} \; \mathsf{mol}^{-1} K^{-1})$ 

K - 2 Cai IIIOI II

A. 163.7 cal

B. Zero

C. 1381.1 cal

D. 9 litre atm

# **Answer: B**



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**29.** The heat of formation of methane at constant pressure is 18500 Kcal at 25°C. What would be the heat of reaction at constant volume?

(Given:  $C(s)+2H_2+CH_4+18500\,$  kcal,  $Q_p=18500\,$ cals, R = 2 kcal,

 $Q_p$  = Heat of reaction at constant pressure)

A. 19096 kcal

B. 18798 kcal

C. 18202 kcal

D. 17904 kcal



**Answer: A** 

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**30.** The work done (in erg) for the reversible expansion of one mole of an ideal gas from a volume of 10 Litres to 20

Litres at 25°C is:

A.  $2.303 imes 298 imes 0.082 \log$  2

B.  $-298 imes 10^7 imes 8.314 imes 2.303 \log$  2

 $\mathrm{C.}-2.303\times298\times0.082\log\,0.5$ 

D.  $2.303 imes 298 imes 2 \log$  2

#### **Answer: B**

