

## Daily Problem Practice-II (CHEMICAL-THERMODYNAMICS)

1. An ideal gas expands in volume from $1 \times 10^{-3} \mathrm{~m}^{3}$ to $1 \times 10^{-2} \mathrm{~m}^{3}$ at 300 K temp against a constant pressure $\mathrm{P}=1 \times 10^{5}$ $\mathrm{Nm}^{-2}$. The work done is -
(a) -900 J
(b) -900 kJ
(c) 270 kJ
(d) +900 kJ
2. Two moles of an ideal gas are expended isothermally and reversibly from 1 L to 10 L at 300 k . The enthalpy change (in kJ ) for process is
(a) +11.4
(b) -11.4
(c) 0
(d) +4.8
3. For the process $\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ [1bar, 373 k$] \rightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$ [1bar, 373 k$]$ the correct set of thermodynamic parameter is -
(a) $\Delta \mathrm{G}=0, \Delta \mathrm{~S}=+\mathrm{ve}$
(b) $\Delta \mathrm{G}=+\mathrm{ve}, \Delta \mathrm{S}=0$
(c) $\Delta \mathrm{G}=0, \Delta \mathrm{~S}=-\mathrm{ve}$
(d) $\Delta \mathrm{G}=-\mathrm{ve}, \Delta \mathrm{S}=+\mathrm{ve}$
4. For a reaction to occur spontaneously:
(a) $(\Delta \mathrm{H}-\mathrm{T} \Delta \mathrm{S})$ must be negative
(b) $(\Delta \mathrm{H}+\mathrm{T} \Delta \mathrm{S})$ must be negative
(c) $(\Delta \mathrm{H})$ must be negative
(d) $(\Delta \mathrm{S})$ must be negative
5. The change in entropy for the following transformations is respectively (+ indicates increases, - indicates decreases and 0 indicates no change)
(i) $\mathrm{SO}_{2} \mathrm{Cl}_{2(\mathrm{~g})} \rightarrow \mathrm{SO}_{2(\mathrm{~g})}+\mathrm{Cl}_{2(\mathrm{~g})}$
(ii) $\mathrm{nCH}_{2}=\mathrm{CH}_{2(\mathrm{~g})} \rightarrow\left[\mathrm{CH}_{2}-\mathrm{CH}_{2(\mathrm{~s})}\right]_{\mathrm{n}}$
(iii) $\mathrm{I}_{2(\mathrm{~s})} \rightarrow \mathrm{I}_{2(\mathrm{v})}$
(iv) Adiabatic reversible expansion an ideal gas
(A) $+,-, 0,+$
(B),,$+- \mathbf{0 , 0}$
(C),,,-++ 0
(D),,,+-+ 0
6. Which of the following relation is correct-
(a) $\Delta \mathrm{G}^{\circ}=\mathrm{RT} \operatorname{lnk}$
(b) $\Delta \mathrm{G}^{\circ}=\mathrm{RT} \log \mathrm{K}$
(c) $\mathrm{K}=\mathrm{e}^{-\Delta \mathrm{G}^{\circ} / 2.303 \mathrm{RT}}$
(d) $\mathrm{K}=10^{-4 \mathrm{G}^{\circ} 2.303 \mathrm{RT}}$
7. $\Delta \mathrm{G}^{\circ}$ for the reaction $\mathrm{A}+\mathrm{B} \rightleftharpoons \mathrm{C}+\mathrm{D}$ is equal to -4.606 kcal . The value of equilibrium constant for the reaction is-
(a) 100
(b) 10
(c) 2
(d) 0.01
8. Which of the following PLOT is correct for relation between $\log K$ and $1 / \mathrm{T}$ for exothermic thermodynamic process
(a) $\log \mathrm{k}$

(b) $\log \mathrm{k}$

(c)

1/T
(d) $\log \mathrm{k}$

9. Which of the following conditions is favourable for the feasibility of a reaction -
(a) $\Delta H=-v e, T \Delta S=+v e$
(b) $\Delta \mathrm{H}=-\mathrm{ve}, \mathrm{T} \Delta \mathrm{S}=+\mathrm{ve}, \mathrm{T} \Delta \mathrm{S}<\Delta \mathrm{H}$
(c) $\Delta \mathrm{H}=+\mathrm{ve}, \mathrm{T} \Delta \mathrm{S}=+\mathrm{ve}, \mathrm{T} \Delta \mathrm{S}<\Delta \mathrm{H}$
(d) $\Delta \mathrm{H}=+\mathrm{ve}, \mathrm{T} \Delta \mathrm{S}+\mathrm{ve}, \mathrm{T} \Delta \mathrm{S}>\Delta \mathrm{H}$
10. The value of $\Delta \mathrm{G}$ for the reaction $n x \rightarrow \mathrm{mB}$, at 700 K if $\Delta \mathrm{H}=-113 \mathrm{~kJ} \mathrm{~mol}^{-1}$ and $\mathrm{DS}=145 \mathrm{JK}^{-1} \mathrm{~mol}^{-1}$
(a) $+11.50 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(b) -11.50 kJmol
(c) $+22.80 \mathrm{~kJ} \mathrm{~mol}^{-1}$
(d) $-30.57 \mathrm{~kJ} \mathrm{~mol}^{-1}$
11. 

$\left(\frac{\partial G}{\partial P}\right)_{T}=$
(a) V
(b) S
(c) -S
(d) $-V$
12. Among the following, the system that would require the highest amount of thermal energy to bring its temperature to $80^{\circ} \mathrm{C}$ is
(a) 400 g of water at $40^{\circ} \mathrm{C}$
(b) 200 g of water at $20^{\circ} \mathrm{C}$
(c) 3000 g of water at $50^{\circ} \mathrm{C}$
(d) 600 g of water at $30^{\circ} \mathrm{C}$
13. Among the following, the reaction that is accompanied by a decrease in the entropy is
(a) $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})$
(b) $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}(\mathrm{~s})+6 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 6 \mathrm{CO}_{2}(\mathrm{~g})+6 \mathrm{H}_{2} \mathrm{O}(l)$
(c) $\mathrm{PCl}_{5}(\mathrm{~s}) \rightarrow \mathrm{PCl}_{3}(l)+\mathrm{Cl} 2(\mathrm{~g})$
(d) $2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \rightarrow 2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})$
14. A thermodynamic process is endothermic and takes place with increase in entropy
(a) is always non-spontaneous
(b) is always spontaneous
(c) spontaneous at high temperature
(d) spontaneous at low temperature

## RAJSI CHEMISTRY POINT <br> For JRF/NET, GATE, DRDO, BARC, IIT-JAM, RPSC and M.Sc. in CHEMISTRY

15. The plot that describes a Carnot cycle is
(a)

(b)

(c)

(d)

16. Which of the following relation is INCORRECT?
(a) $\left(\frac{\partial A}{\partial T}\right)_{V}=-S$
(b) $\left[\frac{\partial A}{\partial V}\right]_{T}=-P$
(c) $\left[\frac{\partial G}{\partial T}\right]_{P}=-S$
(d) $\left[\frac{\partial A}{\partial T}\right]_{T}=P$
17. For a chemical reaction, $\Delta \mathrm{H}^{\circ}=-38.3 \mathrm{~kJ} \& \Delta \mathrm{~S}^{\circ}=-113 \mathrm{JK}^{-1} \mathrm{~mol}^{-1}$. This reaction is -
(a) Spontaneous at all temperature
(b) Non-spontaneous at all temperature.
(c) Spontaneous at temperature above 340 K
(d) Spontaneous at temperature above 340 K
18. The change in entropy when 1 mole of an ideal gas is compressed to $1 / 4$ th of its initial volume and simultaneously heated to twice of its initial temperature is:
(a) $\left[\mathrm{C}_{\mathrm{v}}-\mathrm{R}\right] \ln 2$
(b) $\left[\mathrm{C}_{\mathrm{v}}-2 \mathrm{R}\right] \ln 2$
(c) $\left[C_{v}+2 R\right] \ln 2$
(d) $\left[\mathrm{C}_{\mathrm{v}}-\mathrm{R}\right] \ln 4$
19. A Carnot engine operating between $27^{\circ} \mathrm{C}$ and $127^{\circ} \mathrm{C}$ has efficiency equal to
(a) $21 \%$
(b) $22 \%$
(c) $24 \%$
(d) $25 \%$
20. $T$ - $S$ diagram for a Carnot's cycle is
(a) Rectangle
(b) circle
(c) ellipse
(d) Parabolic
21. The difference in entropy $(\Delta \mathrm{S})$ between a state of volume $V_{i}$ and a state of volume $V_{f}$ (temperature and number of molecules remaining constant) is equal to
(a) $n R \log \frac{V_{f}}{V_{i}}$
(b) $n R \log \frac{V_{i}}{V_{f}}$
(c) $n R^{2} \log \frac{V_{f}}{V_{i}}$
(d) $n R^{2} \log \frac{V_{i}}{V_{f}}$
22. The entropy of an isolated system
(a) Remains unchanged in any process
(b) Remains unchanged or increase during any process
(c) Remains unchanged in an irreversible process
(d) The entropy of non-isolated systems always increase
23. In one of the Maxwell's relations $\left(\frac{\partial S}{\partial p}\right)$ equals
(a) $\left(\frac{\partial V}{\partial T}\right)_{V}$
(b) $-\left(\frac{\partial V}{\partial T}\right)_{p}$
(c) $-\left(\frac{\partial T}{\partial V}\right)_{S}$
(d) $\left(\frac{\partial p}{\partial T}\right)_{V}$
24. Which of the following Maxwell's equation is NOT correct?
(a) $\left(\frac{\partial S}{\partial V}\right)_{T}=\left(\frac{\partial V}{\partial T}\right)_{p}$
(b) $\left(\frac{\partial T}{\partial V}\right)=-\left(\frac{\partial p}{\partial S}\right)_{V}$
(c) $\left(\frac{\partial V}{\partial p}\right)_{S}=\left(\frac{\partial T}{\partial S}\right)$
(d) $\left(\frac{\partial T}{\partial p}\right)_{S}=\left(\frac{\partial V}{\partial S}\right)_{p}$
25. The enthalpy of vaporization of a liquid is $30 \mathrm{~kJ} \mathrm{~mol}^{-1}$ and entropy of vaporization is $75 \mathrm{~J} \mathrm{~mol}^{-1} \mathrm{~K}^{-1}$. The boiling point of the liquid at 1 atm is:
(a) 250 K
(b) 400 K
(c) 450 K
(d) 600 K
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| 11. | 12. | 13. | 14. | 15. | 16. | 17. | 18. | 19. | 20. |
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| 21. | 22. | 23. | 24. | 25. |  |  |  |  |  |
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