## GOVERNMENT OF TAMILNADU HIGHER SECONDARY FIRST YEAR CHEMISTRY

## Unit

7

		THERMODYNAMICS		
Choose th	ie best answer			
The amount of	of heat exchanged with the	ne surrounding at co	onstant pressure is given by the quantity	
a) $\Delta E$	b) <b>Д</b> Н	c) $\Delta S$	d) $\Delta G$	
All the natura	ally occurring processes	proceed spontaneou	usly in a direction which leads to	
a) decrease in	entropy	b) increase in enthalpy		
c) increase in	free energy	d) decrease in free energy		
In an adiabat	ic process, which of the f	following is true?		
a) q = w	<b>b)</b> $q = 0$	c) $\Delta E = q$	d) $P \Delta V = 0$	
In a reversible	e process, the change in	entropy of the unive	erse is	
a) > 0	b) $\geq 0$	c) < 0	d) = 0	
In an adiabat	ic expansion of an ideal g	gas		
a) $\mathbf{w} = -\Delta \mathbf{u}$	b) $w = \Delta u + \Delta H$	c) $\Delta \mathbf{u} = 0$	$\mathbf{d)} \mathbf{w} = 0$	
The intensive	property among the qua	antities below is		
a) mass	b) volume	c) enthalpy	d) mass volume	
-	expands from the volume $\times$ 10 <sup>5</sup> Nm <sup>-2</sup> . The work do		$\times10^{-2}~\text{m}^3$ at 300 K against a constant	
a) - 900 J	b) 900 kJ	c) 270 kJ	d) – 900 kJ	
a) positive The heat of f	bustion is always we b) negative c) ze formation of CO and on of carbon monoxide	CO2 are - 26.4	itive or negative kCal and – 94 kCal, respectively.	

a) +26.4 kcal b) -67.6 kcal c) -120.6 kcal d) +52.8 kcal

10. C(diamond)  $\rightarrow$  C(graphite),  $\Delta H = -ve$ , this indicates that

b) graphite has more energy than diamond d) stability cannot be predicted 11. The enthalpies of formation of $Al_2O_3$ and $Cr_2O_3$ are $-$ 1596 kJ and $-$ 1134 kJ, respectively.						
$\Delta H$ for the reaction $2Al + Cr_2O_3 \rightarrow 2Cr + Al_2O_3$ is						
a) – 1365 kJ	b) 2730 kJ	c) - 2730 l	d) – 462 kJ			
12. Which of the following is not a thermodynamic function ?						
a) internal ene	ergy b) enthalpy	c) entropy	d) frictional energ	У		
13. If one mole of ammonia and one mole of hydrogen chloride are mixed in a closed container to form ammonium chloride gas, then						
a) $\Delta H > \Delta U$	b) ΔH - ΔU	= 0 c) ΔH + Δ	$U=0$ d) $\Delta H < \Delta U$			
14. Change in internal energy, when 4 kJ of work is done on the system and 1 kJ of heat is given out by the system is						
a) +1 kJ	b) - 5 kJ	c) +3 kJ	d) – 3 kJ			
15. The work done by the liberated gas when 55.85 g of iron (molar mass 55.85 g mol $^{-1}$ ) reacts with hydrochloric acid in an open beaker at 25 $^{0}$ C						
a) – 2.48 kJ	b) – 2.22 kJ	c) + 2.22 kJ	d) + 2.48 kJ			
16. The value of $\Delta H$ for cooling 2 moles of an ideal monatomic gas from 125° C to 25° C at constant pressure will be $\left[\text{given C}_p = \frac{5}{2} R\right]$						
a) - 250 R	b) – 500 R	c) 500 R	d) + 250 R			
17. Given that $C(g) + O_2(g) \rightarrow CO_2(g) \Delta H^0 = -a kJ$ ; $2 CO(g) + O_2(g) \rightarrow 2CO_2(g) \Delta H^0 = -b kJ$ ; Calculate the $\Delta H^0$ for the reaction $C(g) + \frac{1}{2}O_2(g) \rightarrow CO(g)$						
a) $\frac{b+2a}{2}$	b) 2a - b	c) $\frac{2a-b}{2}$	d) $\frac{b-2a}{2}$			
18. When 15.68 litres of a gas mixture of methane and propane are fully combusted at $0^{\circ}$ C and 1 atmosphere, 32 litres of oxygen at the same temperature and pressure are consumed. The amount of heat of released from this combustion in kJ is $(\Delta H_{\rm C}~({\rm CH_4}) = -890~{\rm kJ~mol^{-1}}$ and $\Delta H_{\rm C}~({\rm C_3H_8}) = -2220~{\rm kJ~mol^{-1}})$						
a) – 889 kJ	b) – 1390 kJ	c) – 3180 kJ	d) – 632.68 kJ			

c) both are equally stable

a) graphite is more stable than diamond

	tion energy of methane a ond dissociation energy o		mol <sup>-1</sup> and 620 kJ mol <sup>-1</sup> respec-
a) 170 kJ mol <sup>-1</sup>	b) 50 kJ mol <sup>-1</sup>	c) 80 kJ mol <sup>-1</sup>	d) 220 kJ mol <sup>-1</sup>
20. The correct therm Phase - I)	odynamic conditions for	the spontaneous react	tion at all temperature is (NEET
a) $\Delta H < 0$ and $\Delta S$	> 0		
b) $\Delta H < 0$ and $\Delta S$	< 0		
c) $\Delta H > 0$ and $\Delta S$	= 0		
d) $\Delta H > 0$ and $\Delta S$	> 0		
21. The temperature o	f the system, decreases in	an	
a) Isothermal expa	nsion b) Isother	mal Compression	c) adiabatic expansion
d) adiabatic comp	ression		
22. In an isothermal r	eversible compression of a	an ideal gas the sign o	f q, $\Delta S$ and w are respectively
a) +, -, -	b) -, +, -	c) +, -, +	d) -, -, +
23. Molar heat of vap boiling point of the	Section of the sectio	8 kJ mol <sup>-1</sup> . If the entre	ppy change is 16 J mol <sup>-1</sup> K <sup>-1</sup> , the
a) 323 K	b) 27 <sup>0</sup> C	c) 164 K	d) 0.3 K
24. ΔS is expected to	be maximum for the reac	tion	
a) Ca(S) + ½ O <sub>2</sub> (s	$g) \rightarrow CaO(S)$		
b) $C(S) + O_2(g)$	→ CO <sub>2</sub> (g)		
c) $N_2(g) + O_2(g)$	→ 2NO(g)		
d) $CaCO_3(S) \rightarrow C$	$aO(S) + CO_2(g)$		
	and $\Delta S$ for a reaction are e which the reaction will	1 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	$I^{-1}$ and 100 JK <sup>-1</sup> mol <sup>-1</sup> . Then the
a) 300 K	b) 30 K	c) 100 K	d) 20 <sup>0</sup> C