

- 1 Which group of matters the particles are IONS?
- A Polystyrene, Sulphur, Grafite
  - B Zink, Mercury, Sodium chloride
  - C Water, Ethanol, Benzene
  - D Argon, Helium, Diamond
- 2 Which particles are NOT bonded by secondary forces?
- A Water molecules  $\text{H}_2\text{O}$  in ice.
  - B Helium atoms He in gaseous helium.
  - C Nitrogen and oxygen atoms in  $\text{NO}_3^-$  ions.
  - D Molecules of  $\text{CH}_2=\text{CH}_2$ .
- 3 What is the nature of particles in sulphur?
- A S atoms.
  - B  $\text{S}_2$  molecules.
  - C Sulphide ions  $\text{S}^{2-}$ .
  - D  $\text{S}_8$  molecules.
- 4 Which of the following is NOT a primary bond?
- A Hydrogen bond
  - B Elektrovalent bond
  - C Covalent bond
  - D Metallic bond
- 5 From state of matter point of view, what is the feature that make the difference in solid, liquid, and gaseous states of water?
- A Mass of the particles.
  - B Inter-particle distance.
  - C Density of the particles.
  - D The strength of the inter-particle forces.
- 6 In a 0.5 L closed container, 0.6 g of nitrogen gas was sealed. What is the plausible figure to represent the ratio between the total volume of nitrogen molecules to the volume of the container.
- A 1:1
  - B 1:10
  - C 1:100
  - D 1:10 000

- 7 One mole of each of the following matters are contained in identical containers. Which container is the heaviest?
- A Chlorine gas container
  - B Hydrogen gas container
  - C Nitrogen gas container
  - D Helium gas container
- 8 A gas is quoted in mass (or weight). Which of the parameter of the gas that you firstly need in order to estimate its pressure?
- A Temperature
  - B Molar mass
  - C Volume
  - D Universal gas constant
- 9 For an amount of gas at a constant temperature, data on the variation of pressure  $P$  and volume  $V$  were obtained. If the data were plot as  $PV$  versus  $V$  what would you expect the characteristic of the plot?
- A A straight line of positive gradient passing through the origin.
  - B A straight line of negative gradient passing through the origin.
  - C A straight line parallel to the horizontal axis.
  - D A straight line of positive gradient and positive intercept (on vertical axis).
- 10 Which of these properties of gas is not a parameter in the gas state equation?
- A Colour
  - B Molar mass
  - C Volume
  - D Pressure
- 11 What is the kinetic energy of 1 mol of a gas in 1 L volume under 1 atm pressure?
- A 0.0152 J
  - B 0.15 J
  - C 1.5 J
  - D 152 J
- 12 In every collision between any two particles in an ideal gas,
- A Activation energy is achieved.
  - B The energy of the particles before and after the collision are the same.
  - C The velocity of the particles after the collision is lower.
  - D Some of the energy of the particles before the collision is converted into heat after the collision which gives rise to the temperature of the gas.

- 13 What did van der Waals say about the pressure of a real gas in comparison if the gas were ideal?
- A The pressure of a real gas is not affected by the interaction among particles of the gas.
  - B The pressure of a real gas is dependant only on the energy transfer in the collision of the particle with the wall of the vessel.
  - C The pressure of a real gas is lower than the ideal gas due to the interaction among particles of the gas.
  - D The pressure of a real gas is proportional to the interaction forces among the particles of the gas.
- 14 Which of the following is NOT an expression or a statement that formulate the Dalton's Law of partial pressure of mixing gases?
- A  $P = P_A + P_B + P_X$ , when gases A, B, and X are mixed.
  - B The total pressure of mixture of gases A, B and X is the sum of the individual partial pressure of the gases.
  - C  $P_A = x_A P$  where  $x_A$  is the mol fraction of gas A in a mixture, and P is the total pressure.
  - D The expression  $P = P_A + P_B + P_X$ , occurs when gases A, B, and X are miscible.
- 15 12 mol of CO gas under a pressure of 2900 kPa at 298 K was observed to have a volume of 10 dm<sup>3</sup>. Obviously this gas does not behave like an ideal gas. Why? [R = 8.314 JKmol<sup>-1</sup>]
- A The gas is under a very high pressure.
  - B The particle density is too high.
  - C There are too many particles of the gas to behave ideally.
  - D The data do not fit the  $PV = nRT$  relationship.

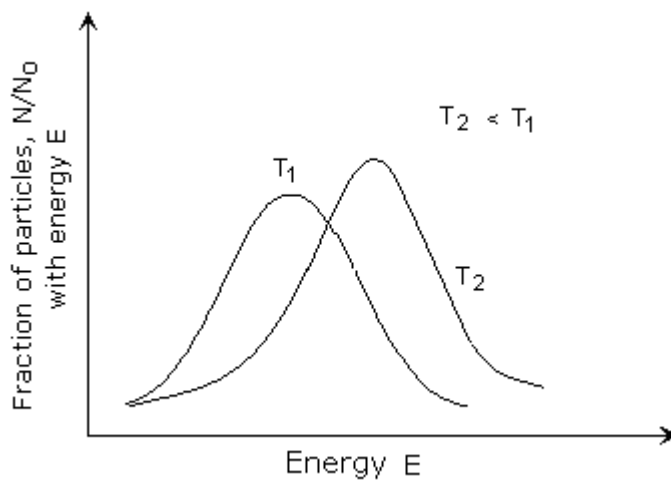


Figure 1

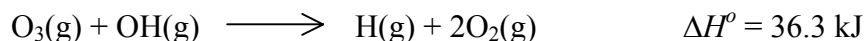
- 16** Figure 1 shows the distribution of energies of particles in a sample of a gas at two different temperature  $T_1$  and  $T_2$ , where  $T_1$  is higher than  $T_2$ . What is NOT right about the plot?
- A The fraction of particles at peak energy for  $T_1$  distribution should be lower than that of  $T_2$  distribution.
  - B The peak energy for  $T_1$  distribution should be at a higher temperature than that of  $T_2$  distribution.
  - C The area under the curve for  $T_2$  distribution should be loer than that of  $T_1$  distribution.
  - D  $T_2$  distribution curve should be broader than that of  $T_1$  distribution.
- 17** n-Octane  $C_8H_{18}$  has a lower vapour pressure than n-hexane  $C_6H_{14}$  although both have very similar molecular nature. What is the possible explanation?
- A Hexane molecules are more spherical than octane molecules.
  - B The heavier octane molecules have less kinetic energy.
  - C Hexane molecules repel each other stronger than that of octane molecules.
  - D Octane has a higher molecular mass, thus has higher van der Waals forces among the molecules.
- 18** The lowest level of sense of smell of  $H_2S$  in human is 0.15 ppb (part per billion). This level MOST PLAUSIBLY means,
- A In every hundred billions (100 000 000 000) of particles in the air, 15 particles are  $H_2S$ .
  - B In every ten billions (10 000 000 000) mol of air, 15 mol are  $H_2S$ .
  - C In every billion (1 000 000 000) g of air, 15 g are  $H_2S$ .
  - D In every billion (1 000 000 000)  $cm^3$  of air, 15  $cm^3$  are  $H_2S$ .
- 19** On average, 3 mg of helium is found in every ton (1000 kg) of earth crust. In term of ppm (part per million), it is,
- A 0.003 ppm
  - B 3.00 ppm
  - C 3000 ppm
  - D 3 000 000 ppm
- 20** The nitrogen content in the air is 75 percent by mass. In term of ppm (part per million), it is,
- A 750 ppm
  - B 7500 ppm
  - C 75 000 ppm
  - D 750 000 ppm

- 21 An  $x\%$  aqueous solution of a solute of  $M \text{ g mol}^{-1}$  molar mass has a density of  $d \text{ g mL}^{-1}$ . What is the *molarity* of the solution?
- A  $10d\left(\frac{M}{x}\right)$   
B  $\left(\frac{10M}{dx}\right)$   
C  $10d\left(\frac{x}{M}\right)$   
D  $\left(\frac{dM}{10x}\right)$
- 22 A glucose ( $\text{C}_6\text{H}_{12}\text{O}_6 = 180 \text{ g mol}^{-1}$ ) solution is  $0.429 \text{ M}$  in concentration with a density of  $1.16 \text{ g mL}^{-1}$ , when expressed in *molality* it is,
- A  $0.20 \text{ m}$   
B  $0.33 \text{ m}$   
C  $0.40 \text{ m}$   
D  $0.396 \text{ m}$
- 23 A solution of a substance with a molar mass  $M \text{ g mol}^{-1}$  is  $m$  molal in concentration and  $d \text{ g mL}^{-1}$  density. The concentration when expressed in *molarity* is,
- A  $\frac{1000md}{1000 + mM}$   
B  $\frac{1000 + md}{1000mM}$   
C  $\frac{md}{mM}$   
D  $\frac{1000md}{mM}$
- 24 A solution of a substance with a molar mass  $M \text{ g mol}^{-1}$  is  $y\%$  in concentration. The *molality* of the solution is,
- A  $\frac{1000y}{100M - y}$   
B  $\frac{1000\left(\frac{y}{M}\right)}{100 - y}$   
C  $\frac{1000y}{M - 100y}$   
D  $\frac{\left(\frac{y}{M}\right)}{100 - y}$

- 25 A solution of phosphoric acid with a molar mass  $98 \text{ g mol}^{-1}$  is 35.4% in concentration. The *molality* of the acid is,
- A 3.62 *m*
  - B 5.59 *m*
  - C 6.71 *m*
  - D 7.26 *m*
- 26 The widely used saline in hospital is a 0.9% solution of sodium chloride in distilled water. Assuming its density is  $1 \text{ g mL}^{-1}$ , what is the molarity of the solution? (Na = 23, Cl = 35.5)
- A 0.17 M
  - B 0.15 M
  - C 0.0015 M
  - D 0.017 M
- 27 MOLAL solution is said to be “thermally-stable” because,
- A Its concentration in molal (*m*) does not change when the temperature of the solution changes.
  - B Molal solution does not dissociate by a temperature change.
  - C Temperature does not affect the solubility of a solute.
  - D Thermal stability of a solution depends on its concentration.
- 28 The following phenomena are colligative properties, EXCEPT,
- A Freezing point depression of a solution
  - B Boiling point elevation of a solution
  - C Osmotic pressure
  - D Dialysis
- 29 Freezing point depression of a solution depends on the physical characteristics of the solute dissolve in it. That characteristic is,
- A the freezing point of the solute
  - B the phase of the solute
  - C the molecular structure of the solute
  - D the number of particles of the solute.
- 30 Ebullioscopic constant of cyclohexane (boiling point,  $80^\circ\text{C}$ ) is  $3 \text{ K kg mol}^{-1}$  ( $\text{K m}^{-1}$ ). A solution X was made by dissolving 10 grams of X (of relative molar mass of 100) in 1 kg of cyclohexane. Solution Y was made by dissolving Y (of relative molar mass of 1000) in 1 kg of cyclohexane. What would be the mass of Y so that the solution Y boils at the same temperature as that of solution X?
- A 1 g
  - B 10 g
  - C 50 g
  - D 100 g

- 31 An amount of  $n$  mol of a solid P was dissolved in a solvent Q at a temperature  $T$  kelvin to give a  $V$  volume of an ideal solution. In terms of these data, this solution shows an osmotic pressure  $\pi$  equals to, [ $R$ , universal gas constant]
- A  $\frac{RT}{nV}$
- B  $\frac{RT}{V} \ln n$
- C  $\frac{nRT}{V}$
- D  $-\frac{nRT}{V}$
- 32 For a change (chemical or physical change) that had taken place in a system,
- A The heat absorbed  $\Delta Q$  by the system is independent of the path along which the change had taken place.
- B The work done  $\Delta W$  by the system is independent of the path along which the change had taken place.
- C The change in internal energy  $\Delta E$  is independent of the path along which the change had taken place.
- D The sum of heat change  $\Delta Q$  and the accompanying work  $\Delta W$  is independent of the path along which the change had taken place.
- 33 During an endothermic chemical reaction,
- A A system becomes warmer, and additional heat is gained from surroundings
- B A system becomes warmer, and the chemical substances undergo an increase in potential energy
- C A system becomes cooler, and the chemical substances undergo an increase in potential energy
- D A system becomes cooler, and the chemical substances undergo a decrease in potential energy
- 34 For a chemical reaction taking place at a constant pressure. Which one of the following is NOT true?
- A  $\Delta E = E_f - E_i$
- B  $\Delta E = E_{\text{product}} - E_{\text{reactants}}$
- C  $\Delta E = q + w$
- D  $\Delta E = \Delta H + P\Delta V$

- 35 For the reaction,  $A_2(g) + 2BC(g) \longrightarrow B_2(s) + 2AC(g)$  taking place in an isolated system, the enthalpy of the product is lower than that of the reactants. Which one of the following statements is true for the system?
- A The potential energy of the system decreases as the reactants are converted to products
  - B The kinetic energy of the system decreases as the reactants are converted to products
  - C The total energy of the system decreases as the reactants are converted to products
  - D The total mass of the system are not changes
- 36 At a constant-volume system, the change in internal energy of the system is
- A  $\Delta E = q_v$
  - B  $\Delta E = C_v$
  - C  $\Delta E = \Delta H$
  - D  $\Delta E = 0$
- 37 A boundary that does permit energy transfer as heat is called
- A adiabatic
  - B exothermic
  - C diathermic
  - D isolated
- 38 Which of the following statements is not true for the internal energy?
- A The internal energy is the total energy of a system
  - B The internal energy is a state function
  - C The internal energy is measured in Joule (J)
  - D If no work is done on the process, the internal energy is equal to zero.
- 39 Use the thermochemical equation

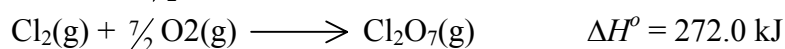
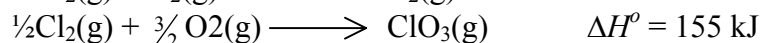
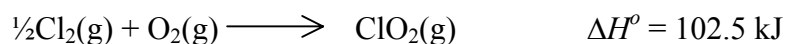


What is  $\Delta H^\circ$  for the reaction of 1.0 g of  $O_3(g)$ ?

- A 0.57 kJ
- B 0.76 kJ
- C 18.15 kJ
- D 36.30 kJ



40 Use the following thermochemical equations:



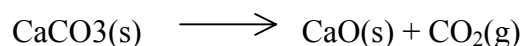
to determine  $\Delta H^\circ$  for  $3\text{ClO}_3(\text{g}) \longrightarrow \text{Cl}_2\text{O}_7(\text{g}) + \text{ClO}_2(\text{g})$

- A -90.5 kJ
- B 90.5 kJ
- C 219.5 kJ
- D 439.0 kJ

**Table 1** The molar enthalpy of substances

Substance	$\Delta_f H^\circ$ (kJmol <sup>-1</sup> )
CaO(s)	-635.6
CaCO <sub>3</sub> (s)	-1206.9
C <sub>2</sub> H <sub>2</sub> (g)	226.6
CO <sub>2</sub> (g)	-393.5
H <sub>2</sub> O(l)	-285.80
O <sub>2</sub> (g)	0

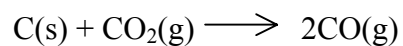
41 Using the data in **Table 1**, calculate the heat of decomposition in the reaction,



- A 42.66 kJ
  - B 1243.58 kJ
  - C 3588.62 kJ
  - D -1243.58 kJ
- 42 Calculate the heat of combustion of acetylene, CH≡CH from the standard enthalpies of formation listed in **Table 1**.
- A 0.0 kJ
  - B -905.9 kJ
  - C -2598.8 kJ
  - D -9059.0 kJ
- 43 For a chemical reaction,  $\Delta H$  is  $< 0$  and  $\Delta S$  is  $< 0$  also. This meant that
- A The reaction must be spontaneous regardless of temperature
  - B The reaction must be spontaneous at high temperature
  - C The reaction may or may not be spontaneous, but spontaneity is favoured at low temperatures
  - D We cannot make any conclusion about spontaneity from the limited information presented.

- 44 Of the species listed below, which one should possess the highest standard entropy ( $S^\circ$ )?
- A  $\text{CH}_4(\text{g})$
  - B  $\text{COCl}_2(\text{g})$
  - C  $\text{C}_2\text{H}_4(\text{g})$
  - D  $\text{C}_4\text{H}_{10}(\text{g})$
- 45 The criterion for a spontaneous chemical reaction is
- A  $\Delta E > 0$
  - B  $\Delta G < 0$
  - C  $\Delta H > 0$
  - D  $\Delta S < 0$
- 46 Which one of the following property in thermodynamics is not a state function
- A enthalpy
  - B entropy
  - C internal energy
  - D work
- 47 Iron undergoes solid-solid transition at  $911^\circ\text{C}$  as it is heated from  $25^\circ\text{C}$  to  $1000^\circ\text{C}$ . Given  $\Delta_{\text{trs}}H^\circ = 0.9 \text{ kJmol}^{-1}$ , calculate  $\Delta_{\text{trs}}S^\circ$  for the changes?
- A  $0.76 \text{ JK}^{-1}\text{mol}^{-1}$
  - B  $0.92 \text{ JK}^{-1}\text{mol}^{-1}$
  - C  $0.99 \text{ JK}^{-1}\text{mol}^{-1}$
  - D  $10.11 \text{ JK}^{-1}\text{mol}^{-1}$
- 48 Predict the enthalpy of fusion of iron at  $1536^\circ\text{C}$  if  $\Delta_{\text{fus}}S^\circ$  is  $9.2 \text{ JK}^{-1}\text{mol}^{-1}$ .
- A  $5.1 \text{ kJmol}^{-1}$
  - B  $9.2 \text{ kJmol}^{-1}$
  - C  $14.1 \text{ kJmol}^{-1}$
  - D  $16.6 \text{ kJmol}^{-1}$
- 49 Consider the reversible expansion of  $1.0 \text{ mol}$  of oxygen from  $25^\circ\text{C}$  and  $24.8 \text{ L}$  to  $427^\circ\text{C}$  and  $38.8 \text{ L}$ . Assuming ideal gas behaviour, calculate  $\Delta S_{\text{sys}}$  for this process. (For  $\text{O}_2(\text{g})$ ;  $C_{p,m} = 29.355 \text{ JK}^{-1}\text{mol}^{-1}$ ,  $C_{v,m} = 21.041 \text{ JK}^{-1}\text{mol}^{-1}$ )
- A  $17.97 \text{ JK}^{-1}\text{mol}^{-1}$
  - B  $21.68 \text{ JK}^{-1}\text{mol}^{-1}$
  - C  $28.79 \text{ JK}^{-1}\text{mol}^{-1}$
  - D  $43.37 \text{ JK}^{-1}\text{mol}^{-1}$

50 Determine  $\Delta G^\circ$  for the reaction



Given  $\Delta H^\circ = 172.46 \text{ kJ}$  and  $\Delta S^\circ = 175.87 \text{ JK}^{-1}$

- A 52.44 kJ
- B 120.02 kJ
- C 168.06 kJ
- D 172.46 kJ

SELAMAT MAJU JAYA

1	2	3	4	5	6	7	8	9	10
B	C	D	A	B	D	A	B	C	A
11									
D	B	C	D	D	B	D	A	A	D
21									
C	D	A	B	B	B	A	D	D	D
31									
C	D	C	D	A	A	C	D	B	A
41									
A	C	C	D	B	D	A	D	B	B