

## Ideal Gas & Kinetic Theory

### Answers and Explanations

#### 1. D

One mole of an ideal gas occupies 22.4 L at STP. This fact you absolutely must memorize for the MCAT. The temperature at STP is 273 K. This is standard temperature. However, 273 K is not *standard state temperature*. Standard state temperature is not 273 K but 298 K. This may seem ridiculous, but there is a figure of merit to knowing about this difference. Many bench-top measurements are at 298 K not 273 K. Charles' Law tells us that at constant pressure the volume of an ideal gas sample is directly proportional to its temperature, i.e.  $V/T = \text{constant}$ , so at 298 K the volume will be greater than 22.4 L.

$$\frac{V_1}{T_1} = \frac{V_2}{T_2}$$

$$\frac{22.4 \text{ L}}{273 \text{ K}} = \frac{24.4 \text{ L}}{298 \text{ K}}$$

#### 2. A

This question is really asking 'when will a real gas behave most like an ideal gas?' Generally, a gas behaves more like an ideal gas at higher temperature and lower pressure, as the potential energy due to intermolecular forces becomes less significant compared with the particles' kinetic energy, and the size of the molecules becomes less significant compared to the empty space between them.

#### 3. B

To use the ideal gas law, first we need to know how many moles is represented by one million atoms.

$$1 \times 10^6 \text{ particles} \left( \frac{1 \text{ mole}}{6.02 \times 10^{23} \text{ particles}} \right)$$

$$= 0.16 \times 10^{-17} = 1.6 \times 10^{-18} \text{ mole}$$

Now we can determine the pressure. Because the answer choices are spaced numerically, you have plenty of latitude for mental math with these computations.

$$PV = nRT$$

$$P = \frac{nRT}{V}$$

$$P = \frac{(1.6 \times 10^{-18} \text{ mole})(8.3 \text{ J mole}^{-1} \text{ K}^{-1})(2.7 \text{ K})}{1 \text{ m}}$$

$$P = \frac{(1.6 \times 10^{-18} \text{ mole})(8.3 \text{ J mole}^{-1} \text{ K}^{-1})(2.7 \text{ K})}{1 \text{ m}}$$

$$P = 3.6 \times 10^{-17} \text{ Pa}$$

#### 4. A

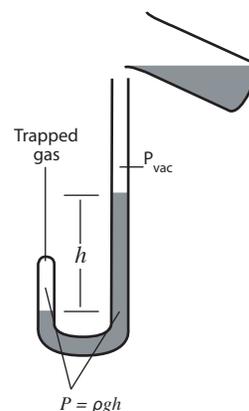
Boyle's Law:

$$P_1 V_1 = P_2 V_2$$

constant temperature

At constant temperature the pressure of an ideal gas sample is inversely proportional to its volume, i.e.  $PV = \text{constant}$ .

The pressure in our trapped gas is directly proportional to the height of the mercury column.



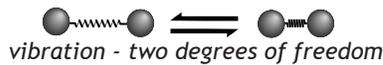
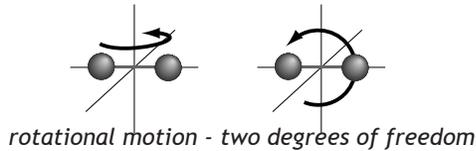
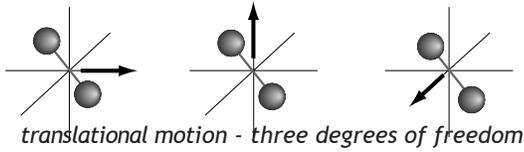
$$P_{\text{g}} V_{\text{g}} = \text{constant} \quad P_{\text{g}} = \rho g h$$

$$\rho g h V_{\text{g}} = \text{constant}$$

$$h V_{\text{g}} = \text{constant}$$

**5. A**

A monatomic gas molecule such as argon possesses only kinetic energy deriving from its linear motion. A diatomic gas molecule, like  $H_2$ , in addition to translational motion, can also rotate and vibrate.



With the ability to store energy in both vibrational and rotational modes, a diatomic gas has more partitions for thermal energy. As a sample of diatomic gas takes in heat, the energy spreads out into all of its degrees of freedom. The other choices besides argon can absorb heat flow into all modes, translational, rotational, and vibrational. For this reason, the molar heat capacity of the other gaseous substances is greater than the molar heat capacity of argon.

**6. C**

Because neon and krypton both exist as monatomic gases, being noble gases, they both possess kinetic energy at the particle level, ie. thermal energy, only in the form of translational kinetic energy. There are no rotational or vibrational modes. If two substances have the same number of places to put kinetic energy at the particle level, they will have the same molar heat capacity. In the case of neon and krypton, their respective constant volume molar heat capacities will be equal and very close to the ideal gas value.

$$C_v = \frac{3}{2} R = 12.5 \frac{J}{mol K}$$

If their molar heat capacities are very close (joules per mole degree Kelvin), then their specific heats will be different (joules per gram degree Kelvin) according to molecular weight.

$$c_{neon} = \left( \frac{12.5 J}{mol K} \right) \left( \frac{mol}{20.2 g} \right)$$

$$c_{kryp} = \left( \frac{12.5 J}{mol K} \right) \left( \frac{mol}{83.8 g} \right)$$

$$c_{neon} : c_{kryp} \approx 4 : 1$$

**7. D**

We can apply the ideal gas model. Helium, a monatomic noble gas, is very near to ideal gas behavior. In a sample of an ideal gas, internal energy is only in the form of thermal energy. The internal energy of an ideal gas is directly proportional the temperature.

$$U = \frac{3}{2} nRT$$

$$3.5 \times 10^6 J = \frac{3}{2} (1 mol) (8.3 \frac{J}{mol K}) T$$

$$T = 2.8 \times 10^5 K$$

**8. A**

Gas B is at a higher temperature. As the temperature of the molecules represented by a Maxwell-Boltzmann distribution increases, the distribution flattens out. rms speed is higher. This corresponds to a greater average kinetic energy per particle, a higher temperature.

**9. A**

The speed at the top of the curve is called the most probable speed because the largest number of molecules have that speed.

**10. D**

The temperature of ideal gas is directly proportional to the translational kinetic energy of the particles. Real gases may have thermal energy in the form of rotational and vibrational kinetic energies as well, so with real gases it's more proper to say that the rela-

relationship is between temperature and kinetic energy per degree of freedom within translational, rotational, and vibrational modes. Nevertheless, whether the gas is ideal or real, temperature increases with the average translational kinetic energy per particle.

$$\frac{1}{2}m\overline{v^2} = \frac{3}{2}kT$$

The particles in gas B are moving at approximately twice the rms speed, so the average kinetic energy per particle is four times greater and so is the temperature.

**11. C**

For this question we assume the distribution represents two gases in thermal equilibrium. In other words, they are the same temperature. At the same temperature, the average translational kinetic energy per particle will be the same, yet the particles of gas B are moving twice as fast on average.

$$\frac{1}{2}m_A\overline{v_A^2} = \frac{1}{2}m_B\overline{v_B^2}$$

If gas B particles are moving twice as fast, to possess the same translational kinetic energy, the mass of gas B particles must be  $\frac{1}{4}$  the mass of gas A particles. The only pair satisfying this condition are  $\text{CH}_4$  (MW 16u) and He (MW 4u).

**12. C**

Neither 'A' nor 'B' is correct because temperature and pressure go up or down together in direct proportionality at constant volume. Both choices 'A' and 'B' would produce no change in the mean free path. Choice 'C' however would increase the mean free path because an increase in volume at constant temperature would lead to a decrease in pressure.

**13. A**

Comparing gases at the same temperature, Graham found experimentally that the rate of effusion of a gas is inversely proportional to the square root of the masses of the particles. This is because effusion rate is proportional to the rms speed of the particles. In

this problem, the temperature of the gas increased. A difference of  $0^\circ\text{C}$  and  $30^\circ\text{C}$  is a difference of 273K and 303K. The Kelvin temperature increased approximately 10%. The average translational kinetic energy of the gas particles is directly proportional to the Kelvin temperature.

$$\frac{1}{2}m\overline{v^2} = \frac{3}{2}kT$$

A 10% increase in Kelvin temperature corresponds to a 10% increase in translational kinetic energy, which, in turn, corresponds to approximately a 3% increase in rms speed ( $\sqrt{10}$ ).

**14. D**

For the buoyant force to be equal and opposite to the weight of the aircraft in both cases (neutral buoyancy), the air inside the hot-air balloon would need to weigh the same as the air inside the vacuum dirigible at 0.5 atm. In other words, it needs to be half as dense as the surrounding air. In a hot air balloon, the air within the balloon is the same pressure as the surrounding air. To possess the same pressure and half the density, the Kelvin temperature needs to be twice as great. You can see this in the ideal gas law.

$$PV = nRT$$

$$\frac{n}{V} = \frac{P}{TR}$$

We are given a temperature for the surroundings of  $27^\circ\text{C}$ , which equals 300K ( $T = T_c + 273.15$ ). Doubling 300K to 600K and converting back to Celsius gives us  $327^\circ\text{C}$ .

**15. A**

This passage has a number of difficult out-of-scope elements. It's important to remember in the exam that a passage like this isn't about foreknowledge of the out-of-scope elements. Ultimately, the questions are going to be about fundamentals and how well you kept your footing. An important figure of merit when you see an unfamiliar equation is to get yourself on speaking terms with it. What is the equation saying? What changes with what?

An elastic modulus indicates how difficult a material is to deform under stress. A material possessing a high elastic modulus has a high rigidity. In the passage we are told that the ratio of elastic modulus to the square of density of diamond is  $E/\rho_s^2 \approx 1.0 \times 10^5 \text{ kg}^{-1}\text{m}^3\text{s}^{-2}$ , but the minimum requirement to prevent buckling for a vacuum dirigible of neutral buoyancy is presented as  $E/\rho_s^2 = 4.5 \times 10^5 \text{ kg}^{-1}\text{m}^3\text{s}^{-2}$ . If a material were developed where the elastic modulus were halved while also quartering density that would increase  $E/\rho_s^2$  eight-fold, so while this material would be less rigid than diamond its lower density (and thus weight) would reduce the stresses it would need to undergo and thus make a viable vacuum dirigible possible.

**16. A**

Neutral buoyancy determines a ratio of shell thickness to radius as a function of the ratio of the density of the surrounding air to the density of the shell material.

$$h/R = \rho_a/(3\rho_s)$$

As the radius increases the shell thickness must increase in direct proportion.

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