iClicker Question:

Which answer represents a list in order of increasing deviation from ideal gas behavior?

Key points of Deviation from Ideal Gas Behavior

- The greater the deviation the less ideal the gas is.
- The deviation from ideal behavior is large at high pressure.
- Low pressure = more ideal (less deviation)
- As temperature increases the deviation from ideal behavior decreases.
- As temperature decreases the deviation increases. (Maximum deviation near the temperature at which the gas becomes a liquid (at point of phase change known as the "critical point").
- Ideal gas particles occupy very little space and have no attraction forces.
- Van der Waals equation used to predict gas behavior:

$$P = (nRT/V - nb) - (n^2a/V^2)$$

A) a. N^2 at 25° C and 1 atm,

b. N² at 25°C and 1 atm,

c. N² at 25°C and 4 atm

Note: $a \rightarrow b$: higher temperature, molecules moving faster so volume increasing.

 $b \rightarrow c$: increase of pressure. (c is less ideal than a.)

Not in correct order!

B) a. CH₄ at 273 K and 1 atm,

b. CH₄ at 273 K and 0.4 atm,

c. CH₄ at 373 K and 1 atm

Note: a is Methane at STP (standard temp. pressure).

 $a \rightarrow b$: lower pressure thus more ideal (particles become farther apart)

 $b \rightarrow c$: higher temp so deviation decreases. (c. is more ideal than b.)

Not in correct order!

C) a. SO₂ at 20°C and 1 atm,

b. SO₂ at 0°C and 1 atm,

c. SO₂ at 0°C and 3 atm

Note: $a \rightarrow b$: Lower temperature (cooling) so volume is also decreasing. Deviation is increasing.

 $b \rightarrow c$: More pressure so larger deviation! Also, volume is decreasing as pressure increases (deviation increases when this occurs). **Correct answer!**

D) H_2O at $200^{\circ}C$ and 1 atm, CO_2 at $200^{\circ}C$ and 1 atm,

Ar at 200°C and 1 atm

Note: a is polar and most likely to deviate (less ideal) b and c are non-polar. c. is most ideal. Not in correct order!

Particles in Phases:

- Gas particles are uniformly distributed and do not display any order to their arrangement. They "bounce" around. Least dense of the 3 common phases.
- Solid particles are most ordered in their arrangement and usually the most dense of the 3 common phases (but there are important exceptions, e.g., water).
- Liquid particles have organization of particles that is in between solid and gas some order but not as much order as in solids. Potential energy is greater in liquids than in solids, but not as great as in gases.
- Temperature is a measure of average kinetic energy. This holds true for solids and liquids as well as for gases.
- Attraction between molecules decrease as temperature increases.
- Increased temperature in a liquid (or solid) = increased vapor pressure. Vapor pressure is the pressure due to the vapor (gas) that forms when some of the particles change phases (liquid to gas).
- Critical point is the condition of pressure and temperature beyond which it is impossible to tell the difference between the liquid and the gas.