The term **polarizability** refers to the tendency of molecules to generate induced electric dipole moments when subjected to an electric field. It originates from the fact that nuclei and electrons are generally not fixed in space and that when molecules are subject to an electric field, the negatively charged electrons and positively charged atomic nuclei are subject to opposite forces and undergo charge separation. Polarizability allows us to better understand the interactions between nonpolar atoms and molecules and other electrically charged species, such as ions or polar molecules with dipole moments.

**Introduction**

Neutral nonpolar species have spherically symmetric arrangements of electrons in their electron clouds. When in the presence of an electric field, their electron clouds can be distorted (Figure \(\PageIndex{1}\)). The ease of this distortion is the polarizability of the atom or molecule. The created distortion of the electron cloud causes the originally nonpolar molecule or atom to acquire a dipole moment. This induced dipole moment is related to the polarizability of the molecule or atom and the strength of the electric field by the following equation:

\[
\mu_{\text{ind}} = \alpha E
\]

where \(\langle E \rangle\) denotes the strength of the electric field and \(\langle \alpha \rangle\) is the polarizability of the atom or molecule with units of \(\text{C m}^2\text{V}^{-1}\).

![Figure \(\PageIndex{1}\): A neutral nonpolar species's electron cloud is distorted by an externally applied electric field. (top) Before the electric field is turned on the electron cloud is isotropically distributed. (bottom) Once the electric field is turned on, an induced dipole moment is generated. If we turn off the electric field, the dipole moment goes away (not shown). (CC BY-NC 4.0; Ümit Kaya via LibreTexts).](image)

In general, polarizability correlates with the interaction between electrons and the nucleus. The amount of electrons in a molecule affects how tight the nuclear charge can control the overall charge distribution. Atoms with fewer electrons will have smaller, denser electron clouds, as there is a strong interaction between the few electrons in the atoms’ orbitals and the positively charged nucleus. There is also less shielding in atoms with fewer electrons contributing to the stronger interaction of the outer electrons and the nucleus. With the electrons held tightly in place in these smaller atoms, these atoms are typically not easily polarized by external electric fields. In contrast, large atoms with many electrons, such as negative ions with excess electrons, are easily polarized. These atoms typically have very diffuse electron clouds and large atomic radii that limit the interaction of their external electrons and the nucleus.
Factors that Influence Polarizability

The relationship between polarizability and the factors of electron density, atomic radii, and molecular orientation is as follows:

1. The greater the number of electrons, the less control the nuclear charge has on charge distribution, and thus the increased polarizability of the atom.
2. The greater the distance of electrons from nuclear charge, the less control the nuclear charge has on the charge distribution, and thus the increased polarizability of the atom.
3. Molecular orientation with respect to an electric field can affect polarizability (labeled Orientation-dependent), except for molecules that are: tetrahedral, octahedral or icosahedral (labeled Orientation-independent). This factor is more important for unsaturated molecules that contain areas of electron-dense regions, such as 2,4-hexadiene. Greatest polarizability in these molecules is achieved when the electric field is applied parallel to the molecule rather than perpendicular to the molecule.

Polarizability Influences Dispersion Forces

The dispersion force is the weakest intermolecular force. It is an attractive force that arises from surrounding temporary dipole moments in nonpolar molecules or species. These temporary dipole moments arise when there are instantaneous deviations in the electron clouds of the nonpolar species. Surrounding molecules are influenced by these temporary dipole moments and a sort of chain reaction results in which subsequent weak, dipole-induced dipole interactions are created (Figure \(\PageIndex{2}\)).

![Figure \(\PageIndex{2}\): A neutral nonpolar species's electron cloud (yellow) can be distorted by a nearby ion (e.g., sodium cation) or polar molecule (e.g., hydrogen chloride) to induce a dipole moment. (CC BY-NC 4.0; Ümit Kaya via LibreTexts).]
These cumulative dipole-induced dipole interactions create attractive dispersion forces. Dispersion forces are the forces that make nonpolar substances condense to liquids and freeze into solids when the temperature is low enough.

Polarizability affects dispersion forces in the following ways:

- As polarizability increases, the dispersion forces also become stronger. Thus, molecules attract one another more strongly and melting and boiling points of covalent substances increase with larger molecular mass.
- Polarizability also affects dispersion forces through the molecular shape of the affected molecules. Elongated molecules have electrons that are easily moved increasing their polarizability and thus strengthening the dispersion forces, (Figure \(\PageIndex{3}\)). In contrast, small, compact, symmetrical molecules are less polarizable resulting in weaker dispersion forces.

![Figure](\PageIndex{3})

\[ V(r) = \frac{-3}{4} \frac{\alpha^2 I}{r^6} \label{2} \]

where

- \(r\) is the distance between the atoms or molecules,
- \(I\) is the first ionization energy of the atom or molecule, and
- \(\alpha\) is the polarizability constant expressed in units of \(m^3\).
This expression of $\alpha$ is related to $\alpha'$ by the following equation:

$$\alpha' = \frac{\alpha}{4 \pi \epsilon_o} \quad \text{(3)}$$

To quantify the interaction between unlike atoms or molecules ($\text{A}$ and $\text{B}$), Equation (2) becomes:

$$V(r) = \frac{-3}{2} \frac{I_A I_B}{I_A + I_B} \frac{\alpha_A \alpha_B}{r^6} \quad \text{(4)}$$

References