

COVALENT COMPOUNDS PROPERTIES

UNDERSTANDING COVALENT COMPOUNDS PROPERTIES: A DEEP DIVE

COVALENT COMPOUNDS PROPERTIES ARE FUNDAMENTAL TO UNDERSTANDING HOW MATTER INTERACTS AND BEHAVES IN THE WORLD AROUND US. UNLIKE IONIC COMPOUNDS, WHICH ARE FORMED BY THE TRANSFER OF ELECTRONS BETWEEN ATOMS, COVALENT COMPOUNDS ARISE FROM THE SHARING OF ELECTRONS. THIS FUNDAMENTAL DIFFERENCE IN BONDING DICTATES A DISTINCT SET OF CHARACTERISTICS, FROM THEIR PHYSICAL STATES TO THEIR ELECTRICAL CONDUCTIVITY. THIS COMPREHENSIVE ARTICLE WILL EXPLORE THE MULTIFACETED NATURE OF COVALENT COMPOUNDS, DELVING INTO THEIR MOLECULAR STRUCTURE, POLARITY, INTERMOLECULAR FORCES, AND HOW THESE FACTORS INFLUENCE THEIR DIVERSE PROPERTIES. WE WILL EXAMINE WHY WATER IS SUCH A UNIQUE SOLVENT, WHY FATS DON'T MIX WITH WATER, AND HOW THE STRUCTURE OF A DIAMOND LEADS TO ITS LEGENDARY HARDNESS. JOIN US AS WE UNLOCK THE SECRETS BEHIND THE BEHAVIOR OF THESE UBIQUITOUS SUBSTANCES.

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WHAT ARE COVALENT COMPOUNDS?

COVALENT COMPOUNDS ARE CHEMICAL SUBSTANCES FORMED WHEN TWO OR MORE NON-METAL ATOMS BOND TOGETHER BY SHARING ELECTRONS. THIS SHARING CREATES STABLE MOLECULES WHERE EACH ATOM ACHIEVES A MORE ENERGETICALLY FAVORABLE ELECTRON CONFIGURATION, TYPICALLY RESEMBLING THAT OF A NOBLE GAS. THINK OF IT AS ATOMS COOPERATING TO FULFILL THEIR ELECTRON "NEEDS" RATHER THAN ONE ATOM COMPLETELY DOMINATING ANOTHER, AS SEEN IN IONIC BONDING. THIS ELECTRON-SHARING MECHANISM IS THE BEDROCK UPON WHICH AN ASTONISHING ARRAY OF CHEMICAL SUBSTANCES IS BUILT, FROM SIMPLE GASES LIKE OXYGEN TO COMPLEX BIOLOGICAL MOLECULES ESSENTIAL FOR LIFE.

THE DRIVING FORCE BEHIND COVALENT BOND FORMATION IS THE DESIRE FOR ATOMS TO ATTAIN A STABLE ELECTRON SHELL. BY SHARING VALENCE ELECTRONS, ATOMS CAN EFFECTIVELY "COUNT" THESE SHARED ELECTRONS TOWARDS THEIR OWN STABLE CONFIGURATION. THIS CREATES A STRONG, DIRECTIONAL BOND THAT DEFINES THE GEOMETRY AND, CONSEQUENTLY, THE PROPERTIES OF THE RESULTING MOLECULE. UNLIKE IONIC BONDS, WHICH ARE TYPICALLY ELECTROSTATIC ATTRACTIONS BETWEEN OPPOSITELY CHARGED IONS AND ARE NON-DIRECTIONAL, COVALENT BONDS ARE LOCALIZED BETWEEN SPECIFIC ATOMS.

THE INFLUENCE OF MOLECULAR STRUCTURE ON COVALENT COMPOUND PROPERTIES

THE ARRANGEMENT OF ATOMS WITHIN A COVALENT MOLECULE, ITS MOLECULAR STRUCTURE, IS A PRIMARY DETERMINANT OF ITS PROPERTIES. THIS INCLUDES THE TYPES OF ATOMS INVOLVED, THE NUMBER OF COVALENT BONDS, AND THE THREE-DIMENSIONAL SHAPE THE MOLECULE ADOPTS. THE GEOMETRY OF A MOLECULE, OFTEN DICTATED BY THE REPULSION BETWEEN ELECTRON PAIRS AROUND THE CENTRAL ATOM (VSEPR THEORY), SIGNIFICANTLY IMPACTS HOW IT INTERACTS WITH OTHER MOLECULES AND WITH EXTERNAL FORCES.

MOLECULAR GEOMETRY AND SHAPE

THE SPATIAL ARRANGEMENT OF ATOMS IN A MOLECULE IS CRUCIAL. FOR INSTANCE, A LINEAR MOLECULE WILL INTERACT DIFFERENTLY WITH ITS SURROUNDINGS THAN A BENT OR TETRAHEDRAL ONE. THIS SHAPE INFLUENCES HOW CLOSELY MOLECULES CAN PACK TOGETHER IN SOLID OR LIQUID STATES, AFFECTING MELTING AND BOILING POINTS. IT ALSO DICTATES THE MOLECULE'S POLARITY, A CONCEPT WE WILL EXPLORE FURTHER. CONSIDER CARBON DIOXIDE (CO_2), WHICH IS LINEAR. WHILE THE INDIVIDUAL CARBON-OXYGEN BONDS ARE POLAR, THE MOLECULE AS A WHOLE IS NONPOLAR DUE TO ITS SYMMETRICAL LINEAR ARRANGEMENT, WHICH CAUSES THE BOND DIPOLES TO CANCEL OUT. THIS HAS SIGNIFICANT IMPLICATIONS FOR ITS SOLUBILITY AND BEHAVIOR.

BOND STRENGTH AND TYPE

THE STRENGTH OF A COVALENT BOND ALSO PLAYS A VITAL ROLE. SINGLE BONDS ARE GENERALLY WEAKER AND LONGER THAN DOUBLE BONDS, WHICH ARE IN TURN WEAKER AND LONGER THAN TRIPLE BONDS. THIS VARIATION IN BOND STRENGTH DIRECTLY AFFECTS THE STABILITY AND REACTIVITY OF THE MOLECULE. STRONGER BONDS REQUIRE MORE ENERGY TO BREAK, LEADING TO HIGHER MELTING AND BOILING POINTS. FOR EXAMPLE, NITROGEN GAS (N_2), WITH ITS STRONG TRIPLE BOND, IS VERY UNREACTIVE AND HAS A LOW BOILING POINT BECAUSE THE INTERMOLECULAR FORCES ARE WEAK.

ALLOTROPES AND THEIR VARYING PROPERTIES

AN INTERESTING ASPECT OF MOLECULAR STRUCTURE IS THE EXISTENCE OF ALLOTROPES. THESE ARE DIFFERENT STRUCTURAL MODIFICATIONS OF THE SAME ELEMENT IN THE SAME PHYSICAL STATE. CARBON, FOR INSTANCE, EXISTS AS DIAMOND AND GRAPHITE. IN DIAMOND, EACH CARBON ATOM IS COVALENTLY BONDED TO FOUR OTHER CARBON ATOMS IN A TETRAHEDRAL ARRANGEMENT, FORMING A RIGID, THREE-DIMENSIONAL NETWORK. THIS STRUCTURE IS RESPONSIBLE FOR DIAMOND'S EXCEPTIONAL HARDNESS AND ELECTRICAL INSULATING PROPERTIES. GRAPHITE, ON THE OTHER HAND, CONSISTS OF LAYERS OF CARBON ATOMS ARRANGED IN HEXAGONAL RINGS. WITHIN EACH LAYER, THE CARBON ATOMS ARE STRONGLY BONDED, BUT THE LAYERS ARE HELD TOGETHER BY WEAKER FORCES. THIS LAYERED STRUCTURE MAKES GRAPHITE SOFT AND A GOOD ELECTRICAL CONDUCTOR, AS ELECTRONS ARE DELOCALIZED WITHIN THE LAYERS.

POLARITY IN COVALENT BONDS AND COMPOUNDS

POLARITY IS A CRITICAL CONCEPT IN UNDERSTANDING COVALENT COMPOUNDS. IT ARISES FROM THE UNEQUAL SHARING OF ELECTRONS BETWEEN BONDED ATOMS, LEADING TO A SEPARATION OF ELECTRICAL CHARGE WITHIN THE MOLECULE. THIS INEQUALITY STEMS FROM THE DIFFERENCE IN ELECTRONEGATIVITY BETWEEN THE BONDED ATOMS. ELECTRONEGATIVITY IS A MEASURE OF AN ATOM'S ATTRACTION FOR SHARED ELECTRONS IN A CHEMICAL BOND.

ELECTRONEGATIVITY DIFFERENCES AND BOND POLARITY

WHEN TWO IDENTICAL ATOMS SHARE ELECTRONS, AS IN AN O_2 MOLECULE, THE SHARING IS PERFECTLY EQUAL, AND THE BOND IS NONPOLAR. HOWEVER, WHEN ATOMS WITH DIFFERENT ELECTRONEGATIVITIES BOND, THE MORE ELECTRONEGATIVE ATOM PULLS THE SHARED ELECTRONS CLOSER TO ITSELF. THIS CREATES A PARTIAL NEGATIVE CHARGE (Δ^-) ON THE MORE ELECTRONEGATIVE ATOM AND A PARTIAL POSITIVE CHARGE (Δ^+) ON THE LESS ELECTRONEGATIVE ATOM. THIS UNEQUAL DISTRIBUTION OF CHARGE CREATES A POLAR COVALENT BOND. FOR EXAMPLE, IN A HYDROGEN CHLORIDE (HCl) MOLECULE, CHLORINE IS MORE ELECTRONEGATIVE THAN HYDROGEN, SO THE ELECTRONS IN THE H-Cl BOND ARE PULLED CLOSER TO CHLORINE, GIVING IT A PARTIAL NEGATIVE CHARGE AND HYDROGEN A PARTIAL POSITIVE CHARGE.

MOLECULAR POLARITY: THE SUM OF BOND DIPOLES

WHILE INDIVIDUAL BONDS CAN BE POLAR, THE OVERALL POLARITY OF A MOLECULE DEPENDS ON ITS GEOMETRY. IF A MOLECULE CONTAINS POLAR BONDS BUT HAS A SYMMETRICAL SHAPE, THE INDIVIDUAL BOND DIPOLES CAN CANCEL EACH OTHER OUT, RESULTING IN A NONPOLAR MOLECULE. WATER (H_2O) IS A CLASSIC EXAMPLE OF A POLAR MOLECULE. THE OXYGEN ATOM IS MORE ELECTRONEGATIVE THAN THE HYDROGEN ATOMS, MAKING THE O-H BONDS POLAR. DUE TO THE BENT SHAPE OF THE WATER MOLECULE, THESE POLARITIES DO NOT CANCEL OUT; INSTEAD, THEY ADD UP TO CREATE A NET DIPOLE MOMENT, MAKING THE OXYGEN SIDE OF THE MOLECULE PARTIALLY NEGATIVE AND THE HYDROGEN SIDES PARTIALLY POSITIVE. THIS POLARITY IS RESPONSIBLE FOR MANY OF WATER'S UNIQUE PROPERTIES.

INTERMOLECULAR FORCES: THE GLUE THAT HOLDS COVALENT MOLECULES TOGETHER

WHILE COVALENT BONDS HOLD ATOMS TOGETHER WITHIN A MOLECULE, WEAKER FORCES, KNOWN AS INTERMOLECULAR FORCES (IMFs), ACT BETWEEN SEPARATE MOLECULES. THESE FORCES ARE CRUCIAL IN DETERMINING THE PHYSICAL PROPERTIES OF COVALENT COMPOUNDS, SUCH AS THEIR MELTING POINTS, BOILING POINTS, AND STATES OF MATTER. THE STRENGTH OF IMFs VARIES GREATLY, AND THEY ARE GENERALLY MUCH WEAKER THAN INTRAMOLECULAR FORCES (COVALENT OR IONIC BONDS).

TYPES OF INTERMOLECULAR FORCES

THERE ARE SEVERAL TYPES OF INTERMOLECULAR FORCES, EACH ARISING FROM DIFFERENT INTERACTIONS:

- **LONDON DISPERSION FORCES (LDFs):** THESE ARE THE WEAKEST IMFs AND ARE PRESENT IN ALL MOLECULES, POLAR AND NONPOLAR. THEY ARISE FROM TEMPORARY FLUCTUATIONS IN ELECTRON DISTRIBUTION WITHIN AN ATOM OR MOLECULE, CREATING INSTANTANEOUS DIPOLES THAT INDUCE TEMPORARY DIPOLES IN NEIGHBORING MOLECULES. LDFs BECOME STRONGER AS THE SIZE AND NUMBER OF ELECTRONS IN A MOLECULE INCREASE, LEADING TO GREATER POLARIZABILITY.
- **DIPOLE-DIPOLE FORCES:** THESE FORCES OCCUR BETWEEN POLAR MOLECULES. THE PARTIALLY POSITIVE END OF ONE MOLECULE IS ATTRACTED TO THE PARTIALLY NEGATIVE END OF ANOTHER. THESE FORCES ARE STRONGER THAN LDFs FOR MOLECULES OF SIMILAR SIZE.
- **HYDROGEN BONDING:** THIS IS A SPECIAL, STRONGER TYPE OF DIPOLE-DIPOLE INTERACTION THAT OCCURS WHEN A HYDROGEN ATOM IS BONDED TO A HIGHLY ELECTRONEGATIVE ATOM (OXYGEN, NITROGEN, OR FLUORINE) AND IS ATTRACTED TO A LONE PAIR OF ELECTRONS ON ANOTHER ELECTRONEGATIVE ATOM IN A NEIGHBORING MOLECULE. HYDROGEN BONDING IS RESPONSIBLE FOR MANY OF WATER'S UNIQUE PROPERTIES, SUCH AS ITS HIGH BOILING POINT AND ITS ABILITY TO ACT AS AN EXCELLENT SOLVENT.

IMPACT ON PHYSICAL PROPERTIES

THE STRENGTH OF INTERMOLECULAR FORCES DIRECTLY CORRELATES WITH THE PHYSICAL PROPERTIES OF COVALENT COMPOUNDS. SUBSTANCES WITH STRONG IMFs, LIKE THOSE WITH EXTENSIVE HYDROGEN BONDING, TEND TO HAVE HIGHER MELTING AND BOILING POINTS BECAUSE MORE ENERGY IS REQUIRED TO OVERCOME THESE ATTRACTIONS AND SEPARATE THE MOLECULES. CONVERSELY, COMPOUNDS WITH ONLY WEAK LDFs, SUCH AS SMALL NONPOLAR MOLECULES, HAVE LOW MELTING AND BOILING POINTS AND ARE OFTEN GASES AT ROOM TEMPERATURE. THIS EXPLAINS WHY METHANE (CH_4), A SMALL NONPOLAR MOLECULE, IS A GAS, WHILE GLUCOSE ($\text{C}_6\text{H}_{12}\text{O}_6$), A LARGER MOLECULE WITH MULTIPLE SITES FOR HYDROGEN BONDING, IS A SOLID.

PHYSICAL PROPERTIES OF COVALENT COMPOUNDS

THE COLLECTIVE IMPACT OF MOLECULAR STRUCTURE, POLARITY, AND INTERMOLECULAR FORCES GIVES RISE TO THE CHARACTERISTIC PHYSICAL PROPERTIES OF COVALENT COMPOUNDS. THESE PROPERTIES ARE WHAT WE OBSERVE AND EXPERIENCE IN OUR DAILY LIVES, FROM THE FEEL OF WAX TO THE BEHAVIOR OF WATER.

MELTING AND BOILING POINTS

COVALENT COMPOUNDS GENERALLY HAVE LOWER MELTING AND BOILING POINTS COMPARED TO IONIC COMPOUNDS. THIS IS PRIMARILY BECAUSE THE INTERMOLECULAR FORCES HOLDING COVALENT MOLECULES TOGETHER ARE WEAKER THAN THE ELECTROSTATIC FORCES IN IONIC LATTICES. BREAKING THESE WEAKER IMFs REQUIRES LESS ENERGY. HOWEVER, THERE ARE EXCEPTIONS. NETWORK COVALENT SOLIDS, LIKE DIAMOND AND SILICON DIOXIDE (SiO_2), HAVE EXTREMELY HIGH MELTING AND BOILING POINTS BECAUSE THEIR ATOMS ARE HELD TOGETHER BY A CONTINUOUS NETWORK OF STRONG COVALENT BONDS, NOT BY DISCRETE INTERMOLECULAR FORCES.

SOLUBILITY

THE PRINCIPLE "LIKE DISSOLVES LIKE" IS A GOOD RULE OF THUMB FOR PREDICTING THE SOLUBILITY OF COVALENT COMPOUNDS. POLAR COVALENT COMPOUNDS TEND TO DISSOLVE WELL IN POLAR SOLVENTS (LIKE WATER), WHILE NONPOLAR COVALENT COMPOUNDS DISSOLVE WELL IN NONPOLAR SOLVENTS (LIKE HEXANE OR OIL). THIS IS BECAUSE SIMILAR INTERMOLECULAR FORCES MUST BE OVERCOME IN BOTH THE SOLUTE AND THE SOLVENT FOR DISSOLUTION TO OCCUR EFFECTIVELY. POLAR SOLUTES CAN FORM FAVORABLE DIPOLE-DIPOLE INTERACTIONS OR HYDROGEN BONDS WITH POLAR SOLVENTS. NONPOLAR SOLUTES CAN INTERACT WITH NONPOLAR SOLVENTS THROUGH LDFs.

STATES OF MATTER

AT ROOM TEMPERATURE AND PRESSURE, COVALENT COMPOUNDS CAN EXIST AS SOLIDS, LIQUIDS, OR GASES. THIS DEPENDS LARGELY ON THEIR MOLECULAR WEIGHT AND THE STRENGTH OF THEIR INTERMOLECULAR FORCES. SMALL, NONPOLAR MOLECULES WITH WEAK LDFs ARE TYPICALLY GASES (E.G., O_2 , N_2 , CH_4). LARGER MOLECULES OR THOSE WITH STRONGER IMFs (DIPOLE-DIPOLE OR HYDROGEN BONDING) ARE MORE LIKELY TO BE LIQUIDS OR SOLIDS (E.G., H_2O , SUGAR, WAX).

ELECTRICAL CONDUCTIVITY OF COVALENT COMPOUNDS

A DEFINING CHARACTERISTIC THAT OFTEN DISTINGUISHES COVALENT COMPOUNDS FROM IONIC COMPOUNDS IS THEIR BEHAVIOR AS ELECTRICAL CONDUCTORS. IN GENERAL, MOST COVALENT COMPOUNDS ARE POOR CONDUCTORS OF ELECTRICITY IN ALL THREE STATES: SOLID, LIQUID, AND GAS. THIS IS BECAUSE ELECTRICAL CONDUCTIVITY REQUIRES THE PRESENCE OF MOBILE CHARGED PARTICLES, EITHER IONS OR FREE ELECTRONS. COVALENT COMPOUNDS, BY THEIR NATURE, CONSIST OF NEUTRAL MOLECULES HELD TOGETHER BY SHARED ELECTRONS. THERE ARE NO FREE-MOVING IONS TO CARRY CHARGE.

WHY THEY DON'T CONDUCT ELECTRICITY

IN IONIC COMPOUNDS, THE STRONG ELECTROSTATIC ATTRACTION BETWEEN POSITIVELY AND NEGATIVELY CHARGED IONS FORMS A CRYSTAL LATTICE. WHEN THESE COMPOUNDS MELT OR DISSOLVE IN WATER, THE IONS BECOME MOBILE AND CAN CARRY AN ELECTRIC CURRENT. COVALENT COMPOUNDS, HOWEVER, REMAIN AS DISCRETE NEUTRAL MOLECULES. EVEN WHEN HEATED TO MELTING OR VAPORIZATION, THE MOLECULES THEMSELVES DO NOT CARRY A NET CHARGE. THE ELECTRONS ARE LOCALIZED

WITHIN THE COVALENT BONDS AND ARE NOT FREE TO MOVE THROUGHOUT THE SUBSTANCE. THEREFORE, THEY CANNOT FACILITATE THE FLOW OF ELECTRICAL CURRENT.

EXCEPTIONS TO THE RULE

WHILE THE GENERALIZATION HOLDS TRUE, THERE ARE A FEW IMPORTANT EXCEPTIONS WHERE COVALENT SUBSTANCES CAN EXHIBIT ELECTRICAL CONDUCTIVITY. GRAPHITE, AS MENTIONED EARLIER, IS A FORM OF CARBON WHERE ELECTRONS ARE DELOCALIZED WITHIN THE LAYERS, ALLOWING IT TO CONDUCT ELECTRICITY. CERTAIN COVALENT COMPOUNDS CAN ALSO BECOME CONDUCTIVE WHEN DISSOLVED IN SPECIFIC SOLVENTS IF THEY UNDERGO A REACTION TO FORM IONS (THOUGH THIS IS LESS COMMON THAN WITH IONIC COMPOUNDS). FURTHERMORE, SOME SPECIALIZED COVALENT MATERIALS, LIKE CERTAIN ORGANIC SEMICONDUCTORS, ARE DESIGNED TO CONDUCT ELECTRICITY UNDER SPECIFIC CONDITIONS.

EXAMPLES OF COVALENT COMPOUNDS AND THEIR PROPERTIES

LOOKING AT REAL-WORLD EXAMPLES HELPS SOLIDIFY OUR UNDERSTANDING OF COVALENT COMPOUND PROPERTIES. THESE SUBSTANCES ARE ALL AROUND US, PLAYING VITAL ROLES IN VARIOUS PROCESSES AND APPLICATIONS.

- **WATER (H₂O):** A POLAR COVALENT MOLECULE WITH EXTENSIVE HYDROGEN BONDING. ITS POLARITY MAKES IT AN EXCELLENT SOLVENT FOR MANY POLAR SUBSTANCES, EARNING IT THE TITLE "UNIVERSAL SOLVENT." ITS HIGH BOILING POINT, HIGH SPECIFIC HEAT CAPACITY, AND SOLID FORM BEING LESS DENSE THAN ITS LIQUID FORM ARE ALL CONSEQUENCES OF STRONG HYDROGEN BONDING.
- **METHANE (CH₄):** THE SIMPLEST HYDROCARBON AND A NONPOLAR COVALENT MOLECULE. IT EXHIBITS ONLY WEAK LONDON DISPERSION FORCES. CONSEQUENTLY, METHANE HAS A VERY LOW BOILING POINT AND IS A GAS AT ROOM TEMPERATURE, COMMONLY USED AS A FUEL.
- **CARBON DIOXIDE (CO₂):** A LINEAR, NONPOLAR MOLECULE DESPITE HAVING POLAR C=O BONDS. ITS NONPOLARITY MEANS IT DOESN'T DISSOLVE WELL IN WATER (THOUGH IT REACTS WITH WATER TO FORM CARBONIC ACID). IT'S A GAS AT ROOM TEMPERATURE AND IS ESSENTIAL FOR PHOTOSYNTHESIS.
- **SUGAR (SUCROSE, C₁₂H₂₂O₁₁):** A POLAR COVALENT MOLECULE WITH NUMEROUS O-H GROUPS, ENABLING SIGNIFICANT HYDROGEN BONDING. THIS LEADS TO ITS SOLID STATE AT ROOM TEMPERATURE AND ITS SOLUBILITY IN WATER.
- **DIAMOND (C):** A NETWORK COVALENT SOLID WHERE CARBON ATOMS FORM A GIANT, RIGID, THREE-DIMENSIONAL LATTICE. THIS STRUCTURE RESULTS IN EXTREME HARDNESS, HIGH MELTING POINT, AND ELECTRICAL INSULATION.
- **GRAPHITE (C):** ANOTHER ALLOTROPE OF CARBON, CONSISTING OF LAYERED STRUCTURES. THE WEAK FORCES BETWEEN LAYERS ALLOW FOR SLIPPERINESS AND MAKE IT A CONDUCTOR OF ELECTRICITY AND HEAT.

FACTORS AFFECTING COVALENT COMPOUND PROPERTIES

WHILE THE FUNDAMENTAL PRINCIPLES OF COVALENT BONDING DICTATE GENERAL TRENDS, SEVERAL FACTORS CAN INFLUENCE THE SPECIFIC PROPERTIES OF A GIVEN COVALENT COMPOUND. UNDERSTANDING THESE NUANCES ALLOWS FOR MORE PRECISE PREDICTIONS AND EXPLANATIONS OF CHEMICAL BEHAVIOR.

MOLECULAR SIZE AND MASS

AS MENTIONED EARLIER, LARGER MOLECULES GENERALLY HAVE STRONGER LONDON DISPERSION FORCES DUE TO HAVING MORE ELECTRONS AND A LARGER ELECTRON CLOUD, WHICH IS MORE EASILY DISTORTED. THIS LEADS TO HIGHER MELTING AND BOILING POINTS FOR LARGER MOLECULES WITHIN A HOMOLOGOUS SERIES OF NONPOLAR COMPOUNDS. FOR EXAMPLE, AS YOU GO UP THE ALKANE SERIES (METHANE, ETHANE, PROPANE, BUTANE, ETC.), BOILING POINTS INCREASE BECAUSE OF INCREASING MOLECULAR SIZE AND LDFs.

PRESENCE OF FUNCTIONAL GROUPS

IN ORGANIC CHEMISTRY, THE PRESENCE OF SPECIFIC FUNCTIONAL GROUPS (E.G., -OH HYDROXYL, -COOH CARBOXYL, -NH₂ AMINO) CAN DRAMATICALLY ALTER THE PROPERTIES OF A MOLECULE. THESE GROUPS OFTEN INTRODUCE POLARITY AND THE ABILITY TO FORM HYDROGEN BONDS, LEADING TO INCREASED SOLUBILITY IN WATER AND HIGHER MELTING/BOILING POINTS COMPARED TO SIMILAR MOLECULES WITHOUT THESE GROUPS. FOR EXAMPLE, ETHANOL (AN ALCOHOL) IS SOLUBLE IN WATER AND HAS A HIGHER BOILING POINT THAN ETHANE (AN ALKANE) DUE TO ITS HYDROXYL GROUP.

CRYSTAL STRUCTURE (FOR SOLIDS)

EVEN FOR COVALENT COMPOUNDS THAT EXIST AS SOLIDS, THEIR SPECIFIC CRYSTALLINE ARRANGEMENT CAN INFLUENCE PROPERTIES LIKE DENSITY AND HARDNESS. FOR NETWORK COVALENT SOLIDS LIKE DIAMOND AND SILICON CARBIDE, THE PRECISE WAY THE ATOMS ARE BONDED IN THREE DIMENSIONS DICTATES THEIR MACROSCOPIC PROPERTIES. SUBTLE DIFFERENCES IN HOW THESE NETWORKS ARE PACKED CAN LEAD TO VARIATIONS IN PHYSICAL CHARACTERISTICS.

THE INTRICATE INTERPLAY OF THESE FACTORS UNDERSCORES THE COMPLEXITY AND DIVERSITY FOUND WITHIN THE REALM OF COVALENT COMPOUNDS. IT'S NOT JUST ABOUT SHARING ELECTRONS; IT'S ABOUT HOW THOSE SHARED ELECTRONS, THE RESULTING MOLECULAR SHAPES, AND THE FORCES BETWEEN MOLECULES ALL CONSPIRE TO CREATE THE UNIQUE PROPERTIES WE OBSERVE.

Q: WHAT IS THE PRIMARY DIFFERENCE IN BONDING BETWEEN COVALENT AND IONIC COMPOUNDS?

A: THE PRIMARY DIFFERENCE LIES IN HOW ATOMS ACHIEVE STABLE ELECTRON CONFIGURATIONS. IN COVALENT COMPOUNDS, ATOMS SHARE ELECTRONS, FORMING DISCRETE MOLECULES. IN IONIC COMPOUNDS, ELECTRONS ARE TRANSFERRED FROM ONE ATOM TO ANOTHER, FORMING CHARGED IONS THAT ARE ATTRACTED TO EACH OTHER IN A CRYSTAL LATTICE.

Q: WHY DO MOST COVALENT COMPOUNDS HAVE LOWER MELTING AND BOILING POINTS THAN IONIC COMPOUNDS?

A: THIS IS BECAUSE THE FORCES HOLDING COVALENT MOLECULES TOGETHER (INTERMOLECULAR FORCES) ARE GENERALLY MUCH WEAKER THAN THE STRONG ELECTROSTATIC ATTRACTIONS BETWEEN IONS IN AN IONIC LATTICE. BREAKING THESE WEAKER INTERMOLECULAR FORCES REQUIRES LESS ENERGY, LEADING TO LOWER MELTING AND BOILING POINTS.

Q: IS WATER A COVALENT OR IONIC COMPOUND, AND WHAT MAKES IT A GOOD SOLVENT?

A: WATER (H₂O) IS A COVALENT COMPOUND. ITS EXCELLENT SOLVENT PROPERTIES ARE DUE TO ITS POLARITY. THE OXYGEN ATOM IS PARTIALLY NEGATIVE AND THE HYDROGEN ATOMS ARE PARTIALLY POSITIVE, ALLOWING WATER MOLECULES TO FORM STRONG HYDROGEN BONDS WITH AND SURROUND THE CHARGED OR POLAR PARTS OF OTHER MOLECULES.

Q: CAN COVALENT COMPOUNDS CONDUCT ELECTRICITY?

A: GENERALLY, NO. MOST COVALENT COMPOUNDS CONSIST OF NEUTRAL MOLECULES AND LACK MOBILE CHARGED PARTICLES (IONS OR FREE ELECTRONS) REQUIRED FOR ELECTRICAL CONDUCTIVITY. EXCEPTIONS INCLUDE GRAPHITE, WHICH HAS DELOCALIZED ELECTRONS, AND SOME SPECIALIZED CONDUCTIVE POLYMERS.

Q: HOW DOES MOLECULAR SHAPE AFFECT THE PROPERTIES OF A COVALENT COMPOUND?

A: MOLECULAR SHAPE INFLUENCES POLARITY. EVEN IF A MOLECULE CONTAINS POLAR BONDS, A SYMMETRICAL SHAPE CAN CAUSE THE BOND POLARITIES TO CANCEL OUT, MAKING THE MOLECULE NONPOLAR. POLARITY, IN TURN, DICTATES HOW MOLECULES INTERACT WITH EACH OTHER AND WITH SOLVENTS, AFFECTING PROPERTIES LIKE SOLUBILITY AND BOILING POINT.

Q: WHAT ARE INTERMOLECULAR FORCES, AND WHY ARE THEY IMPORTANT FOR COVALENT COMPOUNDS?

A: INTERMOLECULAR FORCES ARE THE ATTRACTIVE OR REPULSIVE FORCES THAT EXIST BETWEEN NEIGHBORING MOLECULES. THEY ARE CRUCIAL FOR COVALENT COMPOUNDS BECAUSE THEY DETERMINE THE PHYSICAL PROPERTIES OF THESE SUBSTANCES, SUCH AS THEIR STATE OF MATTER AT A GIVEN TEMPERATURE, MELTING POINT, AND BOILING POINT.

Q: WHY ARE NETWORK COVALENT SOLIDS, LIKE DIAMOND, EXCEPTIONS TO THE LOW MELTING POINT RULE FOR COVALENT COMPOUNDS?

A: NETWORK COVALENT SOLIDS ARE EXCEPTIONS BECAUSE THEIR ATOMS ARE HELD TOGETHER BY A CONTINUOUS, THREE-DIMENSIONAL NETWORK OF STRONG COVALENT BONDS, RATHER THAN BY WEAKER INTERMOLECULAR FORCES. BREAKING THESE STRONG COVALENT BONDS REQUIRES A TREMENDOUS AMOUNT OF ENERGY, RESULTING IN VERY HIGH MELTING POINTS.

Covalent Compounds Properties

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