Addition Reactions Review

This review covers addition reactions from the Alberta Chem 30 organic chemistry unit perspective.

Adding Hydrogen Gas to an Ene

In addition reactions, only the second or third bond in an organic compound is broken.

Consider adding hydrogen gas (H_2) to ethene (C_2H_4) . The second bond in ethene breaks, and one hydrogen atom is added to each side of the double bond.

The product will be two carbons with the four original carbon-hydrogen bonds, plus the two added hydrogen atoms. When the double bond breaks, each carbon gains one extra bonding capacity, and a hydrogen atom can then be added to each side. So, an ene turns into an ane.

$$C_2H_4 + H_2 \to C_2H_6$$

Since hydrogen gas is symmetric from right to left, it does not matter which hydrogen is added to which side; the same product is obtained.

Adding HCl to Ethene and Propene

When adding a molecule that is not symmetric from right to left, you have to consider the potential products that could be formed.

Consider adding HCl to ethene:

- The hydrogen (H) could add to either the left-hand carbon or the right-hand carbon, potentially giving two different products.
- However, in this case, adding the H to either side results in the same molecule: chloroethane.

If you switch the H and Cl around, you still have chloroethane, so you only need to show the molecule once.

Important: Do not draw the same molecule twice if it is, in fact, the same molecule just rotated or flipped.

Adding HCl to Propene

When adding HCl to propene, two different molecules with two different names are obtained. Therefore, both molecules must be drawn.

- 1. First product: Add the H from HCl to the left-hand side and the chlorine to the middle carbon, resulting in 2-chloropropane.
- 2. Second product: Add the chlorine to the terminal carbon and the hydrogen to the middle carbon, resulting in 1-chloropropane.

So, the two possible products are 2-chloropropane and 1-chloropropane.

You also need to balance the chemical reactions. For the chem 30 level, you only need to list the two potential products. In college or university-level chemistry, you will learn which one is more favored (the major product) and which one is the minor. Here, we are listing all possibilities.

$$2 C_3H_6 + 2 HCl \rightarrow C_3H_7Cl (2-chloropropane) + C_3H_7Cl (1-chloropropane)$$

You will need two of the propenes in order to make both of those products. You're going to need two HCl molecules in order to make those two products.

Addition Reaction An addition reaction is a chemical reaction where atoms are added to an unsaturated organic compound (typically containing double or triple bonds), resulting in a saturated compound or a compound with fewer multiple bonds.

Substitution Reactions

This section reviews substitution reactions from the Alberta Chemistry 30 organic chemistry curriculum. In Chem 30, we focus on single substitutions, where one hydrogen atom is replaced by one bromine atom.

The Substitution Process

In a substitution reaction, one of the hydrogen atoms (H) is swapped out for one bromine atom (Br). The goal is to identify all the possible unique products that can result from this single switch. We do not consider scenarios where two or three bromine atoms are added to the same molecule.

For substitution reactions to occur, a **catalyst** is needed, typically in the form of highenergy or ultraviolet radiation. In many cases, **sunlight** can provide enough energy to drive the reaction.

Example: Substituting Propane

Let's consider a three-carbon chain (propane) and examine the products formed when one hydrogen atom is substituted with a bromine atom.

- 1. First Substitution: We start by substituting one of the hydrogen atoms. For example, we can replace one of the hydrogen atoms on the end carbon with a bromine atom.
- 2. Products: This substitution yields **1-bromopropane** and hydrogen bromide (HBr).
- 3. Identifying Equivalent Substitutions: We need to determine which other hydrogen atoms, if substituted, would yield the same product. In this case, substituting any of the hydrogen atoms on the far-left or far-right carbon atoms would result in 1-bromopropane.

Considering Other Possibilities

Now, consider what happens if the hydrogen atom on the middle carbon is replaced in a substitution reaction. This would yield **2-bromopropane**, a different organic product.

Final Products and Balancing

For the given example, only two unique organic products are possible: 1-bromopropane and 2-bromopropane. In addition, two inorganic products result: hydrogen bromide (HBr).

To complete the analysis, the chemical reaction must be balanced. For our specific reaction, two molecules of the original reactant (propane) and two diatomic bromine molecules are needed to carry out the substitution reaction.

Elimination Reactions

Elimination reactions are a type of chemical reaction where atoms or groups of atoms are removed from a molecule, leading to the formation of a double bond (alkene).

Types of Elimination Reactions

There are two main types of elimination reactions:

- Dehydration: Removes water (\$H_20\$).
- Dehydrohalogenation: Removes hydrogen and a halogen (HX).

Both types result in the formation of a double bond where the removed atoms/groups were located.

Dehydration of Alcohols

Dehydration involves the removal of water (\$H_20\$) from an alcohol.

- Requires an acid catalyst (\$H^+\$). Catalysts are written above or below the reaction arrow. Catalysts are not consumed in the reaction.
- The acid catalyst attacks a lone pair on the hydroxy group (OH) of the alcohol.
- This forms a leaving group (\$H_20\$).
- A double bond forms to either the right or left of the carbon that originally had the hydroxy group.
- The hydrogen that is lost replaces the H^+ from the acid.

Example: Dehydration of Butan-2-ol

Starting with butan-2-ol ($CH_3CH(OH)CH_2CH_3$), the A^+ catalyst facilitates the removal of water.

Two possible alkenes can form:

- But-2-ene (\$CH_3CH=CHCH_3\$)
- But-1-ene (\$CH_2=CHCH_2CH_3\$)

The overall reaction:

\$2 \ CH_3CH(OH)CH_2CH_3 \ \xrightarrow{H^+} \ CH_3CH=CHCH_3 \ + \ CH_2=CHCH_2CH_3 \ + \ 2 \ H_2O\$

The key step is to:

- 1. Identify the carbon with the hydroxy group.
- 2. Form a double bond to the adjacent carbon on either side.
- 3. Remove a hydrogen from the carbon where the double bond is formed.

Dehydrohalogenation of Organic Halides

Dehydrohalogenation involves the removal of a hydrogen (\$H\$) and a halogen (\$X\$) from an organic halide.

- Requires basic conditions, typically using hydroxide (\$OH^-\$) as a reactant, not a catalyst.
- The hydroxide helps to remove a hydrogen, which facilitates the removal of the halogen and the formation of a double bond.
- The double bond forms to either the right or left of the carbon that originally had the halogen.

Example: Dehydrohalogenation of 2-Chlorobutane

Starting with 2-chlorobutane (\$CH_3CHClCH_2CH_3\$), the hydroxide reactant facilitates the removal of \$HCl\$.

Two possible alkenes can form:

- But-2-ene (\$CH_3CH=CHCH_3\$)
- But-1-ene (\$CH_2=CHCH_2CH_3\$)

The overall reaction:

\$2 \ CH_3CHClCH_2CH_3 \ + \ 2 \ OH^- \ \rightarrow \ CH_3CH=CHCH_3 \ + \ CH_2=CHCH_2CH_3 \ + \ 2 \ CL^-\$

In this reaction:

- The halogen leaves as a halide ion (\$Cl^-\$).
- The hydrogen removed by the hydroxide forms water (\$H_2O\$).

Spectator Ions

Sometimes, the hydroxide reactant is shown with a spectator ion, such as sodium (\$Na\$) in sodium hydroxide (\$NaOH\$).

Spectator Ion: An ion that is present in the reaction mixture but does not participate in the actual chemical reaction.

The sodium ion does not affect the reaction and can be ignored for balancing purposes.

Example with the spectator ion:

\$2 \ CH_3CHClCH_2CH_3 \ + \ 2 \ NaOH \ \rightarrow \ CH_3CH=CHCH_3 \ + \ CH_2=CHCH_2CH_3 \ + \ 2 \ NaCl\$

Complete Combustion with Generic Formulas

Combustion is the rapid reaction between a substance with an oxidant, usually oxygen, to produce heat and light. Complete combustion of a hydrocarbon yields carbon dioxide and water vapor.

Homegap

To balance a combustion reaction, you must know the hydrogen:carbon ratio. The general formula for alkanes is C_nH_{2n+2} .

Example: Combustion of 2,2-dimethyl Decane

2,2-dimethyl Decane contains:

- Dec- Stem with 10 carbons.
- Two methyl groups, each with one carbon.

Total carbon count = 10 + 2(1) = 12, hence C_{12} .

Since it is an alkane, we use 2n+2 to find the hydrogen count where n = 12:

 $$2(12) + 2 = 26$, hence H_{26}.$

Our hydrocarbon is \$C_{12}H_{26}\$.

The balanced combustion reaction is:

 $C_{12}H_{26} + \frac{37}{2}O_{2(g)} \$ Combustion Balancing Tips

When balancing combustion equations, remember that oxygen needs to be in pairs, or an even number. If you have an odd number of oxygen atoms because of an odd number of hydrogen atoms (e.g., 26 hydrogen atoms make 13 pairs, leading to an odd number of oxygen atoms), double the fuel to achieve an even number.

For example, if you have $C_{12}H_{26}$, doubling the fuel gives $2 * C_{12}H_{26}$. This results in 26 pairs of hydrogen atoms or 26 water molecules. With an even number, balancing becomes more straightforward.

In this case, \$2*12 = 24\$ carbons will yield 24 carbon dioxide molecules. The oxygen atoms on the product side include 48 from carbon dioxide (\$24*2\$) and 26 from water, totaling 74. Since 74 is divisible by 2 (37 pairs), you avoid fractional coefficients for oxygen. Without doubling, you would have obtained a fractional answer of \$37/2\$ for oxygen.

Incomplete Combustion

Incomplete combustion occurs when there is a lack of oxygen (\$O_2\$ is the limiting reagent). In this scenario, not every carbon atom can fully saturate with two oxygen atoms.

Products of Incomplete Combustion:

- Carbon Dioxide (\$CO_2\$): A greenhouse gas.
- Carbon Monoxide (\$CO\$): A toxic gas that is colorless and odorless. It binds to hemoglobin in the blood, preventing oxygen binding and leading to suffocation.
- Carbon (Soot): Black soot, which is harmful to breathe in. High concentrations in the air can lead to poor air quality.
- Water Vapor (\$H_20\$): Not an environmental concern.

Review Guide Structure

Consider creating a review guide using the following structure for organic chemistry families:

Family	Characteristic/Functional Group	Example	Name of Example
Aromatics			
Cyclos			
Organic Halides			
Alcohols			
Carboxylic Acids			
Esters			

For reactions, a review can summarize the number of products and examples:

Reaction Type	Number of Products	Example(s)
Addition	One or Two	Symmetric addition, asymmetric addition
Esterification		Reaction between a carboxylate and an alcohol to make an ester
Elimination		Elimination of water, elimination of H and a halogen
Substitution		
window.MathJax =		
{		

```
tex: {
  inlineMath: [['$', '$'], ['\\(', '\\)']],
  displayMath: [['$$', '$$'], ['\\[', '\\]']]
}
```

};

Elimination Reactions

Dehydration of Alcohols

- Involves the elimination of water (H_2O) from an alcohol to form an alkene.
- Requires an acid catalyst (e.g., H_2SO_4) for the reaction to occur. The acid catalyst helps in the removal of the hydroxyl group (OH) as water, which is a good leaving group.
- The acid catalyst is not consumed in the reaction; it is regenerated.

Catalyst: A substance that increases the rate of a chemical reaction wit

- A double bond forms between the carbon that was attached to the OH group and an adjacent carbon, with the concurrent elimination of a hydrogen atom.
- Generalized reaction:

$$Alcohol \xrightarrow{Acid \ Catalyst} Alkene + H_2O$$

Dehydrohalogenation of Organic Halides

- Involves the elimination of a hydrogen halide (e.g., HCl, HBr) from an organic halide to form an alkene.
- Requires basic conditions. The base is a reactant and is consumed in the reaction.
- The base (e.g., NaOH) assists in the removal of a proton (H^+), facilitating the formation of a double bond.
- Multiple possible alkene products may form, depending on the structure of the organic halide and the position of the halogen atom.
- Generalized reaction:

$$Organic\ Halide \xrightarrow{Base} Alkene + Halide\ Salt + Water$$

Factors Affecting Product Formation

When multiple alkene products are possible in dehydrohalogenation, consider the following:

- Position of the Halogen Atom: The location of the halogen on the carbon chain determines the possible positions for the double bond.
- Possible Products: Different alkenes can be formed depending on which adjacent carbon loses a hydrogen atom.
- Example Scenario: With a four-carbon chain organic halide (e.g. Chlorobutane), the double bond could form between carbon 1 and 2 (but-1-ene) or between carbon 2 and 3 (but-2-ene) after HCl elimination.

Reaction Conditions Comparison

Feature	Alcohol Dehydration	Organic Halide Dehydrohalogenation
Reaction Type	Elimination	Elimination
Reactant	Alcohol	Organic Halide
Reagent	Acid Catalyst	Strong Base
Role of Reagent	Catalyst (regenerated)	Reactant (consumed)
Byproducts	Water	Water and Halide Salt
Product	Alkene	Alkene

Ester Formation: Esterification

Esters are the last family of organic compounds we will study, focusing on their formation and naming conventions.

Ester Reaction Overview

Ester formation involves a reaction between an organic compound derivative (carboxylic acid) and another hydrocarbon derivative (alcohol).

- The reaction is unique due to the combination of these two types of reactants.
- Alcohols can undergo elimination reactions with acid catalysts, but esterification involves a carboxylic acid (containing a \$-C=00H\$ group) reacting with an alcohol (containing a \$-0H\$ group).
- The products are an ester and water. It's crucial to remember the water byproduct.
- A strong, concentrated acid is required as a catalyst, such as sulfuric acid (\$H_2SO_4\$).

Example: Propanoic Acid + Methanol

Reacting propanoic acid with methanol in the presence of a strong acid yields an ester and water.

The carboxylic acid is too weak to act as a catalyst itself.

Ester Formation Mechanism

While the detailed mechanism is beyond the scope of this course, understanding the basic connection is essential.

- 1. The alcohol's oxygen (with lone pairs of electrons) connects to the carbon in the carboxylic acid's carbonyl group (\$C=0\$).
- 2. This carbon is electron-deficient due to the polar carbon-oxygen bond.

Electronegativity: Oxygen has a higher electronegativity than carbon, making the carbonyl carbon electron-poor and susceptible to nucleophilic attack by the alcohol oxygen.

- 3. During the reaction, the \$-OH\$ from the carboxylic acid and a hydrogen from the alcohol are eliminated, forming water.
- 4. The catalyst facilitates this process.

What is an Ester?

Esters are a distinct organic family characterized by a carbon bonded to an oxygen, which is then bonded to another carbon within the main chain. The general structure is \$R-C00-R'\$, where \$R\$ and \$R'\$ are hydrocarbon groups.

Naming Esters

Esters derive their names from the alcohol and carboxylic acid reactants. IUPAC nomenclature dictates the following:

- 1. Alcohol Portion: Named first, using the alkyl name (methyl, ethyl, propyl, butyl, etc.).
- 2. Carboxylic Acid Portion: Named second, using the alkane name derived from the acid's stem. Drop the "-e" and add "-oate".

Example: Ethanoic Acid + Propanol

Reacting ethanoic acid with propanol in the presence of a strong acid catalyst will produce an ester and water.

• The oxygen from propanol connects to the carbonyl carbon of ethanoic acid.

Naming the Product

- 1. Identify the alcohol and carboxylic acid fragments in the ester. The C=0 is a key indicator of the acid portion.
- 2. Name the alcohol portion:
 - Propanol (3 carbons) becomes "propyl".
- 3. Name the carboxylic acid portion:
 - Ethanoic acid (2 carbons) becomes "ethanoate".
- 4. Combine the names:
 - The ester's name is propyl ethanoate.

The "-oate" suffix signifies the ester linkage (C=0-0-).

Practice

Let's test your understanding with an example:

Draw the structure of ethyl methanoate.

- 1. Identify the alcohol and acid portions:
 - Ethyl (alcohol portion)
 - Methanoate (acid portion)
- 2. Draw the structure:
 - Start with the ester linkage (\$C=0-0-\$).
 - Add the ethyl group (\$CH 2CH 3\$) to one side of the oxygen.
 - Add the methanoyl group (\$H\$) to the carbonyl carbon.

Reaction Types

Esterification is an important reaction to remember. Below is a table showing all the reactions studied in this course.

Reaction	Description
Addition	Two or more reactants combine to form a single product.
Substitution	An atom or group in a molecule is replaced by another atom or group.
Elimination	A molecule loses atoms or groups, often forming a double bond. Includes E1 and E2 mechanisms.
Combustion	A rapid reaction between a substance with an oxidant, usually oxygen, to produce heat and light. Can be complete (producing \$C0_2\$ and \$H_20\$) or incomplete (producing \$C0\$, C , and H_2O `).
Esterification	A reaction between a carboxylic acid and an alcohol to form an ester and water, catalyzed by a strong acid.

Naming Esters

Let's walk through naming esters with a few more examples. Remember, the key is to identify the alcohol and acid components.

Methyl Ethanoate Example

Consider methyl ethanoate. Here's how it's structured:

- Two line segments lead into the oxygen without any carbons.
- The carbons are located at three key points.

Butanoic Acid Example

Let's synthesize an ester from butanoic acid and ethanol.

- Butanoic acid has four carbons and a double-bonded oxygen (OOH).
- An acid catalyst is needed (strong, concentrated acid).
- The oxygen (*O*) from the alcohol's hydroxyl group connects to the carbonyl carbon.

The resulting ester will have the following structure:

- Four carbons with a double bond.
- An oxygen atom (O) from the alcohol.
- Two more carbons.

Don't forget the water molecule (H_2O) that is produced as a byproduct, important for balancing the equation.

Naming the Ester

- 1. Alcohol portion first: In this case, it's ethyl (two carbons).
- 2. Acid portion second: Butanoic acid becomes butanoate (four carbons). The "ane" is dropped and replaced with "oate".

So, the ester's name is ethyl butanoate. We are focusing on non-branched examples.

Benzoic Acid Example

One more example using benzoic acid.

- Benzoic acid: An aromatic hydrocarbon derivative with a carboxyl group (COOH) attached to a seventh carbon, not directly to the ring.
- The connection sequence remains the same.

The arrow goes from the alcohol oxygen to the carbon of the $C=\mathcal{O}$ bond.

Reaction

The alcohol (e.g., methanol) reacts with the benzoic acid:

- 1. One carbon from the alcohol.
- 2. The black oxygen (O) stays.
- 3. Connects to a blue carbon with a double-bonded oxygen (C=O).
- 4. The blue oxygen (O) from the acid goes away as part of the water molecule (H_2O).
- 5. The aromatic ring remains.

Don't forget the water!

Naming

The alcohol part (methyl) comes first. Benzoic acid becomes benzoate by dropping the "oic acid" and adding "oate," resulting in methyl benzoate.

Ester Properties and Uses

Esters are formed through esterification.

Esterification: A chemical reaction between an alcohol and an acid to form an ester and water.

- Methanol and benzoic acid have hydrogen bonding, making them soluble in water.
- Esters exhibit dipole-dipole forces but lack hydrogen bonding, so they aren't as soluble.

Esters tend to have distinct smells. They may smell like bananas, peaches, or have a more pungent odor like nail polish remover.

Hydrocarbon Cracking

One last topic.

Hydrocarbon Cracking: Breaking long hydrocarbons into smaller, more valuable pieces by heating them under controlled conditions.

- Atom count must match on both sides of the reaction.
- For example, $C_{20}H_{42}$ can be broken into $C_9H_{20}+C_{11}H_{22}$

Context

Fractional distillation diagrams often appear on exams. Be aware that hydrocarbon cracking is often listed as a wrong answer choice when identifying fractional distillation.

Esterification Reactions

Esterification reactions involve the reaction between an alcohol and a carboxylic acid to produce an ester and water. This is unique in Alberta Chemistry 30 as it is the only reaction that requires two organic reactants.

Reaction Overview

During the reaction, the oxygen (O) from the alcohol connects to the carbon (C) that is double-bonded to an oxygen (O) – the carbonyl – and a hydroxyl group.

Visually, if ethanol (an alcohol) reacts with methanoic acid, the following occurs:

- 1. The oxygen from the ethanol connects to the carbon in the methanoic acid. The carbon to hydrogen bonds in the ethanol do not change.
- 2. The O-H bond in the alcohol breaks, allowing the alcohol to connect to the carbon from the methanoic acid.
- 3. The hydroxyl group (OH) from the methanoic acid is released.
- 4. The hydroxyl group combines with the hydrogen (H) from the alcohol to form water (H_2O).

 $Alcohol + CarboxylicAcid \rightarrow Ester + Water$

Ester Uses

Esters are prevalent in the food industry due to their pleasant smells and are often synthesized to produce specific scents in products like gum.

Catalyst

To facilitate the reaction, an acid catalyst is required. Sulfuric acid is a common catalyst used in esterification reactions.

Naming Esters

Naming esters involves two parts: the alcohol component and the acid component.

- 1. Alcohol Component: The alcohol part is named first.
 - Identify the number of carbons in the alcohol.
 - Use the corresponding prefix (e.g., "eth-" for two carbons).
 - Add "-yl" to the prefix.
 - Example: If the alcohol has two carbons (ethanol), the first part of the ester's name is "ethyl."
- 2. Acid Component: The acid part is named second.
 - Identify the number of carbons in the acid.
 - Use the corresponding prefix (e.g., "meth-" for one carbon).
 - Drop the "-e" from the alkane name.
 - Add "-oate" to the stem.
 - Example: If the acid has one carbon (methanoic acid), the second part of the ester's name is "methanoate."
- 3. Combining the Names: Combine the names of the alcohol and acid components to form the ester's name.
 - Example: Ethyl methanoate.

Example

For the ester formed from ethanol and methanoic acid:

- The alcohol part (ethanol) has two carbons, so it becomes "ethyl."
- The acid part (methanoic acid) has one carbon, so it becomes "methanoate."
- The ester is named ethyl methanoate.

Hydrocarbon Combustion Reactions

Complete Combustion

- This type of combustion is not new and has been covered in previous science courses.
- A hydrocarbon fuel, such as methane (CH_4) , reacts with oxygen (O_2) .
- Requires sufficient or excess oxygen.
- Produces carbon dioxide gas (CO_2) and water vapor (H_2O) .
- Both carbon dioxide and water vapor are greenhouse gases.
 - Water vapor is often overlooked in discussions about greenhouse gases.
- Example of a balanced equation:

$$CH_4 + 2O_2 \longrightarrow CO_2 + 2H_2O$$

Incomplete Combustion

- Occurs when oxygen is the limiting reagent.
- The ratios of products depend on how limiting the oxygen is.
- Observed in vehicles, such as transport trucks emitting black smoke when going uphill or accelerating, indicating incomplete combustion.
- Balancing incomplete reactions will not be assessed.
- Focus is on the products and their environmental effects.
- The products of incomplete combustion are:
 - Carbon dioxide (CO_2): A greenhouse gas.
 - Carbon monoxide (CO):

A deadly, odorless, and tasteless gas. Carbon monoxide detectors are essential in homes to alert occupants of leaks, commonly from furnaces.

• Pure Carbon (*C*):

Soot that can be seen in fireplaces, barbecues, or emitted from transport truck smokestacks.

• Water Vapor (H_2O)

Comparison of Combustion Types

Feature	Complete Combustion	Incomplete Combustion
Oxygen Supply	Sufficient or excess	Limiting reagent
Products	Carbon dioxide, water vapor	Carbon dioxide, carbon monoxide, pure carbon (soot), water vapor
Unique Products	None	Carbon monoxide, soot
Repeating Products	Carbon dioxide, water vapor	Carbon dioxide, water vapor

Polymerization

In this lesson, we're shifting our perspective to understand polymers, which are ginormous molecules unlike the other families we've studied.

Basic Polymer Terminology

Here's some key terminology for understanding polymers:

- Monomers: These replace the term "reactants." A *mono*mer is one unit that turns into *many* units.
- Polymers: This term replaces "product." Polymers are the result of many monomers joining together.

Polymers often have masses of millions of grams per mole, forming long chains that weave around each other, giving plastics their flexibility. Because of the complexity, we use a shorthand to represent these molecules.

Addition Polymerization

Addition polymerization involves breaking a double bond to create connections.

- The second bond in a double bond is a π bond, which is easily broken.
- Breaking the π bond frees up a connection on each side. Instead of adding small molecules, another monomer connects to each end.
- Each connection frees up another end, allowing continuous growth.

Combining Ethene Monomers

Let's look at what happens when we put two ethene monomers together. The π bond breaks, creating bonding spots, and the carbons at the end of the double bond connect.

You can stop the process of building a polymer by adding a chemical likechlorine that terminates the chain, but that's not covered in this lesson.

Typically, we draw polymers with "n" repeating units, where n can be any whole number. The dashes show the connections coming in and out of the bracket.

Naming Convention

The naming convention for polymers involves adding "poly" in front of the monomer name. For example, the polymer made from ethene is called **polyethene**. The "ene" comes from the double bond that used to exist in the monomer, but there's no double bond in the polymer itself.

Everyday Examples

Recycling codes 2 and 4 are both forms of polyethylene, with 4 being high-density polyethylene HDPE, often used in milk jugs.

Polypropene Example

If we start with propene, we get polypropene.

Here's how it works:

- 1. The double bond breaks.
- 2. The carbon at the end of the double bond connects to another carbon at the end of the double bond.
- 3. The CH_3 group remains unaffected.

This results in a branched polymer structure that is more complex than polyethylene. The repeating pattern is shown "n" times.

Whenever you break the double bond, there are always four connections around

Recycling code 5 is polypropylene, often found in water bottles.

Polystyrene

Using styrene as a monomer, which has a benzene ring, we create polystyrene.

Here's the repeating pattern:

- Two carbons that were part of the double bond
- No double bond anymore
- Three connections are H's
- One connection is a phenol side group

Polystyrene is a very common plastic, often marked with recycling code 6. It can be found in two forms:

- Styrofoam: Air is pumped in during production, making it light and fluffy.
- Hard and Dense: No air is added, resulting in a denser material, like the brown Tim Hortons cups.

Mixing Monomers

Chemists and companies like DuPont are always experimenting with mixing different monomers to create new products with mass market appeal. Many times, these discoveries are accidental.

One example is Scotch tape. The glue was originally intended to be permanent, but it ended up being removable, which proved to be a useful property.

Example: Bromoethene and Tetrachloroethene

If we mix bromoethene and tetrachloroethene together, we get a repeating unit with one of each. If mixed in the right ratio, it will keep alternating. The repeating pattern will be BrH followed by CCl_2 .

Polymerization Reactions

Backwards Testing

Alberta Education has started testing students on polymers by showing them a polymer and asking what monomer created it. Students must understand the bonds well enough to identify the family or draw the structure of the monomers. For example, a question might present a polymer and ask which of the multiple-choice answers shows the correct monomer that makes up the polymer.

Addition Polymerization

In addition polymerization, a double bond is broken to connect monomers.

- The monomer must be an -ene, not an -ane, because the double bond is essential for the reaction.
- It is possible to be asked what two monomers make a given polymer, requiring careful examination of the repeating pattern.

Condensation Polymers

Condensation polymers undergo a dehydration reaction, similar to ester formation, but are categorized differently because the reaction repeats extensively.

• What sets them apart from ester chemistry is all of the monomers have two functional groups, allowing connection on both sides.

Dehydration Reaction: A chemical reaction that involves the removal of a water molecule from a reactant.

Process

- 1. Di-functional acid and di-functional alcohol monomers react.
- 2. An alcohol can react on one side of a carboxylic acid and extend the chain, while another carboxyl group on the other side can also grow the chain to the right.
- 3. The hydroxyl group on the alcohol can grow to the left and right, creating a polymer instead of just an ester.

During esterification, the alcohol connects to a carbon atom, and a hydroxyl group is removed, initiating the connection.

$$C = O$$

The hydroxyl group that comes off connects to the carbon.

Visualizing Polymers

Imagine the process happening repeatedly, growing the polymer chain.

Model Representation

Acid - Alcohol - Acid - Alcohol...

Polyesters and Trademarked Names

In general, anything that is condensation polymerization that is bi-functional, you will make a polyester.

- Polyester is a general term for polymers made this way.
- Nylon is a trademarked name by DuPont for a specific type of polyester.

Byproducts

Two water molecules are produced as byproducts, one from each end of the reaction. Balancing these reactions is typically not required.

Flammability of Polyesters

Polyester materials are highly flammable and should be kept away from heat sources.

Biological Polymers

After discussing synthetic polymers, we can transition to biological polymers.

- Amino acids make up DNA.
- Starches are polymers made out of sugar.

Biological Molecules

Tri-Fatty Acids

- Tri-fatty acids are a type of biological molecule.
- They involve a polyester type of chemistry occurring three times.

Glycerol: A three-carbon alcohol with an -OH group on each carbon, allowing it to form ester linkages through reactions with carboxylic acids. The length of the acid piece can vary.

- The product has three carbons from the alcohol, each with an ester linkage.
- Tri-fatty acid is the polymer, while glycerol and fatty acids are the monomers.

Amino Acids, DNA, and RNA

- Amino acids can be linked together to form dipeptides (two amino acids) or polypeptides (many amino acids).
- There are 20 different amino acids that serve as the building blocks of life.
- Important monomer-polymer pairs to be familiar with:
 - DNA
 - RNA

Carbohydrates and Sugars

Created by **Turbolearn AI**

- When eating sugars, you're consuming carbohydrates and cellulose.
- Monosaccharides are the monomers.
 - Putting two sugars together creates a disaccharide.
 - Putting many together forms polysaccharides.
- Examples of monosaccharides include glucose and fructose.
- Examples of polysaccharides:
 - Carbohydrates that humans can consume.
 - Celluloses (grasses) that cows can break down but humans can't.
 - Complex carbohydrates (like pastas) take longer for our bodies to break down, providing energy more slowly.
- Connect the monomers and polymers:
 - Glucose (monomer) and carbohydrate (polymer).

Cracking

Hydrocarbon Cracking

Created by **Turbolearn Al**

- Cracking is a process of breaking down compounds, crucial for the Alberta energy industry.
- Often, what comes out of the ground isn't what customers want, necessitating the breaking down of compounds.

Cracking: A reaction where a large molecule is broken into two or more smaller molecules using heat and pressure.

- Example: a large molecule $C_{20}H_{42}$ is broken into $C_{11}H_{22}$ (diesel fuel or gasoline) and C_9H_{20} .
- When cracking, you get varying pieces, not uniform results.
- Essential for taking what comes out of the ground and making it sellable, especially in Alberta's oil sands, which contain longer-chained hydrocarbons.
- Cracking is less necessary for light sweet oil from places like Saudi Arabia, which already contains more smaller chains.

Introduction to Polymers

In organic chemistry, we're moving into a section focused on **polymers**. These are large structures, and instead of using the term "reactants," we'll use "monomer" to describe the single unit that builds up into the larger structure.

A monomer is a single unit that can join together with other monomers to form a larger structure called a polymer.

The resulting products are called "polymers," consisting of many repeating units (tens of thousands or even hundreds of thousands). The molar mass of polymers can reach into the millions per mole. Polymers are flexible and are the plastics we use daily.

A polymer is a large molecule made up of many repeating subunits called monomers.

IUPAC Naming Conventions

Monomers are named following standard organic chemistry nomenclature. Polymer IUPAC names indicate the type of polymer but not its exact size.

For example:

• poly(monomer name)

This is a general classification, indicating that many monomers have come together.

Addition Polymerization

Addition polymerization is our first type, where monomers add to each other. Think of it like the addition reactions you've learned, where a double bond breaks. Let's start with two monomers and then expand to many.

Ethene Example

Consider ethene (two carbons with a double bond) as our monomer. In addition chemistry, the second, weaker bond breaks, allowing monomers to link up via carbon-carbon connections.

$$CH_2 = CH_2 + CH_2 = CH_2 \rightarrow -[CH_2 - CH_2 - CH_2 - CH_2] -$$

At each end, another monomer can add. To represent a large number of monomers, we can show a repeating pattern.

If we have ten ethene molecules, the extra bond breaks, and connections form. At each side of the monomer where the double bond used to be, future connections can occur. To show this repeated unit, we use square brackets with lines coming out, indicating repetition n number of times.

$$-[CH_2 - CH_2]_n$$

This polymer's common name is polyethylene. It is used in number two and number four recycling plastic codes, commonly found in milk jugs and containers.

Propene Example

Next, consider **propene** (three carbons with a double bond). We can create **polypropene** from it.

Starting with two propene monomers:

- There are always four substituents around the double bond.
- These substituents will show up in our repeating unit.

When two units come together, the monomer propene makes the polymer polypropene. This is the number three plastic code.

The typical exam representation is one unit with n of those:

$$-[CH_2 - CH(CH_3)]_n -$$

Styrene Example

Styrene is the monomer used to create **polystyrene** (styrofoam). Styrene contains a benzene ring.

When you break the double bonds of styrene and link the units together, you get polystyrene.

Two styrene units connecting:

$$-[CH_2 - CH(C_6H_5)]_n -$$

These are your number six plastics. The density can vary, resulting in styrofoam packing chips or Tim Horton's lids, which are denser forms of polystyrene.

Creative Examples

Let's consider a more creative example with bromine and chlorine atoms. Suppose we have monomers with bromine and chlorine in the molecule, creating an alternating pattern.

If we have two carbons in the repeating unit, with a Cl, H, and H, we can deduce that we need an alkene to have made this polymer. For example, 1-bromo-2-chloroethene.

Condensation Polymerization

Condensation polymers involve the same chemistry as ester reactions, but the monomers have two functional groups that can form ester linkages. This involves two carboxyl groups in the acid and two hydroxyl groups in the alcohol.

You add a large amount of each monomer, and the hydroxyl group in the carboxylic acid connects to the carbon in the alcohol, forming an ester linkage.

In this reaction, water is produced. This reaction occurs at both ends, creating a repeating pattern on each side. This results in a repeating chain of acid-alcohol-acid-alcohol.

Polyesters: Building Blocks and Decoding

Let's explore how to identify the monomers that make up a polyester, working backward from the polymer structure.

Identifying Monomers in Polyesters

- 1. Locate the Oxygen in the Backbone: The "O" in the repeating backbone pattern is key. This helps identify the acid and alcohol pieces that combined to form the ester linkage.
- 2. Find the Carbonyl Group: Look for the carbonyl group (C=O). This is the acid side. The other side connected to the oxygen will be the alcohol side.

Building Back the Monomers

Let's consider a polyester with a repeating unit represented as "rocket" repeating n times.

 $[rocket]_n$

To decode this polyester, we need to identify the acid and alcohol monomers that formed it.

- 1. Acid Component: If the acid portion has two carbons, construct a full carboxyl group on both sides of those two carbons to complete the acid component.
- 2. Alcohol Component: On the alcohol side, determine the number of carbons and ensure you identify the correct number of hydroxyl groups (-OH). Wrong answer choices on a test might have the correct carbon count but an incorrect number of hydroxyl groups.

Example

Imagine our polyester repeating unit has a two-carbon acid component and a one-carbon alcohol component. The monomers would be:

- Acid: A two-carbon diacid (i.e., an acid with carboxyl groups on both ends).
- Alcohol: A one-carbon dialcohol (i.e., an alcohol with two hydroxyl groups).

Key Vocabulary

Monomer: A molecule that can bond to other identical molecules to form a polymer.

Polymer: A substance or material consisting of very large molecules, or macromolecules, composed of many repeating subunits.

Created by **Turbolearn Al**

Polyester: A polymer containing ester functional groups in its main chain.

Carbonyl Group: A functional group composed of a carbon atom double-bonded to an oxygen atom (C=O).

Hydroxyl Group: A functional group consisting of an oxygen atom bonded to a hydrogen atom (-OH).

Polymer Skills

- Predicting polymers from monomers.
- Predicting monomers from polymers (working backward).

No need to worry about drawing these structures for the test. Focus on being able to interpret diagrams, work forward and backward, and name the compounds involved.

Which of '					-		-								_			\vdash	\rightarrow	-	
+ +	the foll	owing m	ionome	rs is	used	to cre	eate p	olyst	yrene	comm	only k	now	n as	styı	ofoc	ım?					
thene																					
																		-			
ropene	Styr	ene is t	he mor	nomer	. usec	to c	reate	polys	tyren	e, whic	h is c	omm	only	kno	wn a	ıs					
tyrene	1 1	ofoam a						• •	,	ļ			1								
1,710.10			 																		
olyethyle	ne																				
Which o	t tha t	allawina	nalum	ore is	form	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\		h 000	donca	ion no	luman	i a a ti.		d.	icina		+02		a h	.,,,,	مطارو
Which δ	i ine i	ollowing	Potyiii	e13 13	10111	ieu ir	ii oug	n con	aensa	юп ро	iyillel	ΙΖάΤΙ	ori, p	oroat	icing	wa	iei	as	αв	ург	bauc
Polyethy	lene																				
В		olystyre	no ic m	oada f	from	cturar	. ma	nomo	rc thr	ough (additio	n no	lume	riza	tion	Th	ic n	roc	000	inve	ماداد
Polystyr		reaking				•						٠.					•				
C								•								1					
Polyeste		olyester ionomer:																			, wr
D	"	onomer.	3 WIIII	100	uncin	Jilai Ç	jioup	5 101	ii esie	IIIINO	iges a	iu pi	ouu	LE W	uiei	us	u L	יקעי	ouu	C 1.	
Polyprop	ene																				
Consider	r the t	wo main	types	of po	olymei	rizatio	n dis	cusse	d: ad	dition	and co	nde	nsati	on.	Addi	tion					
polymer	ization	involves	direc	t linki	ng of	mond	mers	, whi	e con	densat	ion po	lyme	riza	tion	invo	lves	th	e fo	orm	atio	n of
small mo	olecule,	such as	water	r, as a	ı bypr	oduct	. Rec	all w	hich t	pe inv	olves	este	r linl	kage	s.						
polymer	is repr	esented	by th	e stru	ıcture	: -[CH	2-CH	(Cl)]n	Wh	ch mo	nomer	was	use	d to	cre	ate	this	s po	olym	er?	
		ne polym	ner -[C	H2-CH	H(Cl)]r	n- is f	forme	d fro	m chl	proeth	ene (a	lso k	now	n as	viny	l ch	ılor	ide)	. Dı	ırin	g
nloroethe	ine) po	olymeriz	ation,	the do	ouble	bond	in ch	loroe	thene	break	s, allov	wing	the	mor	ome	rs 1	o li	nk	toge	ethe	er.
																		-			
hene																		1 1			
opene																					
ropene																					
ropene tyrene onsider t							'	1.					•								
ropene tyrene onsider t	Focus	on the s	smalles	t repe	eating	g unit	'	1.					•								
ropene tyrene onsider t	Focus	on the s	smalles	t repe	eating	g unit	'	1.					•								
ropene ryrene onsider t	Focus	on the s	smalles	t repe	eating	g unit	'	1.					•								
opene yrene onsider t	Focus	on the s	smalles	t repe	eating	g unit	'	1.					•								
ropene tyrene onsider t	Focus	on the s	smalles	t repe	eating	g unit	'	1.					•								
ropene tyrene onsider t	Focus	on the s	smalles	t repe	eating	g unit	'	1.					•								
thene ropene tyrene onsider t ackbone? ructure	Focus	on the s	smalles	t repe	eating	g unit	'	1.					•								

A			٦.																			1:0	· .									
A chemi		-	_			Ī	. 1								·					_					٠.	ope	rtie	S. V	vnic	n n	non	ome
would b	e most	sui	tabl	e fo	r cr	reat	ing	а	poly	yme	er v	vith	a	rep	eati	ng	uni	t of	-L	CH(CH	3)-C	H2.	n-?								
A																																
Ethene																																
В	Prop	ene	(CH	зсн	=CH	2)	wou	ıld	res	ult	in	a po	olyr	ner	wit	th d	ı re	peo	atin	a u	nit	of ·	-[Cŀ	I (CI	1 3)-	-CH	21n	- a	fter			
Propene																		•		9			•		,		•					
С	Polyi)	,,,,,		α	1110	uo	uD(J.I.G.																				
Styrene																																
D																																
1-bromo	_2_ch	loro	eth <i>a</i>	ne																												
Rememb																									_						- 1)
Consider	the s	truc	tur	e of	ead	ch r	mon	om	er	and	l ho	w i	t w	oul	d co	ntr	⁺ibu	te	to 1	the	rep	eat	ing	un	it o	f tl	ne p	oly	me	r. A	sk	
yourself	: Whic	h of	the	ese	mor	nom	ers	со	nta	ins	a n	netk	ıyl	gro	up	(CH	3)	tha	t co	ould	be	pr	ese	nt i	n t	he	rep	eat	ing	uni	t	
shown?																																
A polyme	er has	the	rep	eat	ing	uni	t -[CH	2-C	H(C	6H	5)]n	ı \	Nho	ıt m	non	ome	r is	s us	sed	to	cre	ate	thi	s p	olyr	ner	, ar	nd w	vha'	t is	the
common										-															·							
A						J 1	•																									
Ethene;	Dalvet	hyla	ne																													
В	i otyet	,.																														
	Dali m																															
Propene:	Polyp	rope																														
C				The	rep	oea [.]	ting	j ur	nit -	-[Cŀ	12-	CH(C6F	[5)]	n- (cor	resp	on	ds	to p	oly	sty	ren	e, w	vhic	h is	cr	eat	ed 1	fror	n t	he
Styrene;	Polys	tyre	ne	mor	nome	er s	styr	ene	2, C	ont	aini	ng	a b	enz	ene	rir	ng (C6F	1 5).													
D																																
1-bromo	-2-chl	oroe	the	ne;	Poly	/(1-	bro	mo	-2-	chl	oro	ethe	ene)																		
Think abo	ut th	o cti	cuct	1120	+ha	ı+ ie	مادد	400		har	770	20.	ina	· .	nci	dor		a+	ma	nor	205	roc		in	20	lvct	vro	na	Day	iou	, +4	
													_								161	163	ulli	3 III	РО	ıysı	yıe	IIE.	REV	/IEV	V 11	ie
examples	от ро	ıyme	ers (ana	mor	ıom	ers	ın	The	S TE	TXS	то	mat	cn	The	re	pea	TINÇ	gui	nit.												
A scienti	st dis	cove	rs a	ı ne	w p	olye	este	r v	vith	а	rep	eat	ing	uni	t co	onto	ainir	ng d	a th	ree	2-C	arbo	n c	icid	СО	mpa	ner	nt d	ınd	a t	wo-	-
carbon a	lcohol	con	npor	ent	. Wł	nich	СО	mb	inat	lion	of	mo	non	ner	s w	oul	d fo	rm	th	is p	oly	este	er?									
A														t pol	vest	er i	s foi	rmed	d fr	om a	dia	cid d	and .	a dia	ılcoh	ol. A	thr	ee-	carbo	on d	iacid	
A three-	-carbo	n di	acid	and	d a	two)-ca	ırba	on d	diale	coh	ol.		-			_		_	_	_			_					des t			
В	94.70												F	orti	on o	f th	e re	pea	ting	unit	in	the 1	olye	ester								
A three-	carbo	n m	2006	ocid	and	ا ما	+1440		arh	on I	mar	امما	coh	ما ،	٨٨	200																
	-cai bo	11 1110	J110C	LIU	uilo	· u	ı wC	<i>)</i> -c(ui Di	U11 1	iior	ισαι	CON	UI. U	∕∪ (<	710	3															
С																	Ĺ					-										
A two-c	arbon	diac	ıd a	ind	a th	ree	e-ca	ırbo	on d	iale	coh	ol.																	Ш			
D																																
A two-c	arbon	mon	oac	id a	nd d	a th	ree	2-C	arb	on I	mor	noal	coh	ol.																		
A polyes	ter re	auir	oc +	wo	fun	ctic	nal	ar	OUT	S 0	n c	ach	m	no	mer	· (a	dia	ciط	an	d 4	dia	ارما	رام	to	on/	ıbla	th	o n	olvr	nor	iza:	tion
' '		•						_	٠.						WY			Sid	uii	<u>. u</u>	ui0		101)	10	SIIC	ع) د		٦ ٢	51711		u	1011
process	ui DOT	n er	us,	cre	uiin	y a	rep	rea	ıınğ	j po	או ופ	ı II.					J															
																													Ш			
			T																													

Which of	the fo		·	•						_				•	•				ma	non	2	c lin	\L	hu	forr	mine	2 05	tor b	ands
Polyester	*											•														mile	, es	iei bi	Jilus
В]		Γ								[• [the	dor	uble t	ond
Polyethyle																											-		10114
A polyme	r form	od f	rom (a dia	cid	ana	4 4	lialca	hal																				
D D		leu i	10111	a aid	ciu	un	u u u	iluicc	71101																				
A polyme	r with	este	r link	anes	s in	th	e ma	in cl	ain																				
71 potymo				ugu	J		<u> </u>	01																					
A new pol	ymar i	e evi	nthas	izad	wit	lh f	lha r	anaa	tino		i+	ſсц	2 (·LL/E	2~11~	. \	M/ha	1+ i	c +1	ر مو آ	מנוי	۸,	nan	2	of 4	tha	mar	omai	usad
to create	1	•		izeu	WII		ille i	epec	ııııç	uii	" -	ĮCH	2-0	,,,,	ינל יי		VVIIC	A1 1	3 11	16 1	.UF	i	IUII	HE	01 1	116	111011	ome	useu
Δ Creare	iiiis pi	Jiyiiii	51 3																										
poly(brome	athan	(0																											
Pora(prouit	Jemen	E																											
poly(ethen	a hror	nida)																											
Polyternen	e DI OI	iliue)																											
poly(1 bro	maath	anal	The	mor	nom	ier	used	is 1	-br	omo	eth	ene	, w	hick	n po	olyn	neri	zes	to	for	m l	ooly	(1-	bro	mo	ethe	ene).	. The	'1-'
poly(1-bro	moern	ene) indi	cates	s th	nat	the	bron	nine	ato	m i	s at	tac	hec	t to	th	e fir	rst	car	bor	ı in	the	et	he	ne r	nole	3Cule	3.	
poly(2-bro	maath	ana)																											
poly(2-bio	illoein	enej																											
																										-			
		l		-													-							- 1.					
A scientis		1 1	-	1 1				and	dide	enti	fies	the	e re	epe	atin	g u	nit (as	-[CI	H2-	CH(CH2	CH	3)]	n `	Wh	at m	ionon	ner
was used	to syr	thes	ize tl	his p	olyı	mer	r?																			_			
A																										-	_	_	
Propene																										-	_		
	he rep		_																										
Butene								neriz	zatio	n,	the	dou	ıble	: bo	nd	bre	aks,	, al	low	ing	bu [.]	tene	: m	ond	me	rs t	o lir	ık	
	ogethe	r, fo	rming	g the	po	olyn	ner.																			_	_		
Ethene																										_			
D																													
Styrene																													
																												_	
A chemist	aims	to sy	nthe	size	ар	oly	mer	simil	ar t	οр	olys	tyr	ene	bu	ıt us	sinc	j a r	mor	nom	er	wit	h ar	ı e	thy	l gr	our) (CH	12CH	3)
attached	to the	carl	on ir	ıstea	ıd o	of a	ben	zene	rin	g (C	6H:	5). \	٧h٥	at v	voul	ld b	e t	he	rep	eat	ing	uni	t o	f t	his	new	/ pol	ymer	?
A			Т	he e	thv	Lar	roup	(CH2	CH:	3) r	enla	res	th	e h	enz.	ene	rin	a (C6F	1 5)	on	the	ca	rho	n a	tom			
-[CH2-CH	CH2CI	(3)]n	1			_	rep				•							9 (.	,	•	0		. 50					
В					9		, i op		.g		ĮŪ.		٠. ١٠	•		٠,].	' '												
-[CH2-CH	(C6H5)]n-																											
С																													
-[CH(CH3)	-CH2]	n-																											
D																													
-[CH2-CH	(Br)]n-																									\neg			
			61 1 6 ±	ne -	رے ع	- عدد الم		ام اما	- +	_	Al. :-	مدد ۱	or -	n		m l-	~ <u>'</u>	- al-	١	- C.L.	.	n+ -	11	o le	ر ہے۔	ta 1	h a a	عامره.	a in
Consider h						·				٠.	*	•																	
the monor								•	eatii	ıg ı	JNIT.	of	The	po	ıym	er.	ıne	n C	ons	iae	r ho	ow 1	ne	et	nyı	gro	Jup	snoul	a
replace th	e ben	∠ene	ring	on 1	ıne	cai	roon.		-								-	-		-		-	-	_		\rightarrow	-	-	

[CH2-CHCI]n- B -[CH2-CH2]n- C -[CHCI-CHCI]n-				1 1 1							
[CH2-CH2]n- ; [CHCl-CHCl]n-											
[CHCl-CHCl]n-											
CH3-CHCl]n-											
nink about addition poly onnect with other monor nit based on the structu	mers. Considei	r what ator	ms will be d					_			atir
Consider a polymer fo	ormed from	the mond	omer 2-ch	loro-1,3-b	outadiene	$(CH_2$	= C	'Cl -			
$CH=CH_2$). If this repeating unit of the r			addition p	oolymeriz	<i>ation</i> , wh	at wou	ıld be	the			
A -[CH2-CCI=CH-CH		•	merization, chloro-1,3-								215
[CH2-CCI-CH-CH			le bond shif		•	_					
P (2010 201 201 201	2]n-										
B -[CH2-CH=CCI-CH C -[CH3-CCI=CH-CH	2]n-										
C -[CH3-CCI=CH-CH D -[CH2-CCI=CH-CH member that addition	3]n- polymerizatio			T							tio
C -[CH3-CCI=CH-CH D -[CH2-CCI=CH-CH member that addition gle bonds to create a rowing the monomers to	3]n- polymerizatio repeating unit o link. Think a	t. In the mo lbout where	onomer 2-c e the chlori	nloro-1,3-l	outadiene, 1	he dou	ble bo	nds v	will br	eak,	
C -[CH3-CCI=CH-CH D -[CH2-CCI=CH-CH member that addition gle bonds to create a rowing the monomers to	3]n- polymerizatio repeating unit o link. Think a	t. In the mo lbout where	onomer 2-c e the chlori	nloro-1,3-l	outadiene, 1	he dou	ble bo	nds v	will br	eak,	
C -[CH3-CCI=CH-CH D -[CH2-CCI=CH-CH member that addition gle bonds to create a rowing the monomers to	3]n- polymerizatio repeating unit o link. Think a	t. In the mo lbout where	onomer 2-c e the chlori	nloro-1,3-l	outadiene, 1	he dou	ble bo	nds v	will br	eak,	
C -[CH3-CCI=CH-CH D -[CH2-CCI=CH-CH member that addition gle bonds to create a rowing the monomers to	3]n- polymerizatio repeating unit o link. Think a	t. In the mo lbout where	onomer 2-c e the chlori	nloro-1,3-l	outadiene, 1	he dou	ble bo	nds v	will br	eak,	
C -[CH3-CCI=CH-CH D -[CH2-CCI=CH-CH member that addition gle bonds to create a rowing the monomers to	3]n- polymerizatio repeating unit o link. Think a	t. In the mo lbout where	onomer 2-c e the chlori	nloro-1,3-l	outadiene, 1	he dou	ble bo	nds v	will br	eak,	
C -[CH3-CCI=CH-CH D -[CH2-CCI=CH-CH member that addition gle bonds to create a rowing the monomers to	3]n- polymerizatio repeating unit o link. Think a	t. In the mo lbout where	onomer 2-c e the chlori	nloro-1,3-l	outadiene, t	he dou	ble bo	nds v	will br	eak,	
C -[CH3-CCI=CH-CH	3]n- polymerizatio repeating unit o link. Think a	t. In the mo lbout where	onomer 2-c e the chlori	nloro-1,3-l	outadiene, t	he dou	ble bo	nds v	will br	eak,	

Which of the following is	NOT a	typica	l cha	arac	teris	tic o	fрo	lyme	rs?													+
A Composed of repeating mo	onomer	units.																				1
B																						+
Formed through addition (or cond	ensatio	n re	actio	ons.																	+
					rs h											r lar	ge :	size	, re	sul	ing	+
ow molar mass compared	to mor	nomers.	fro	m t	he m	any	rep	eatin	g ma	non	ner	unit	s, n	ot lo	w.							+
D Can be flexible, like many	plastic	s																				
A scientist discovers a nev							٠,	'														
two-carbon chain, which kely used to create this			ѕ то	an o															ers	are	e mo	ST
\								er fo				1		T								_
thanoic acid and methan	ol.		4					mpor							•							tes
		لد مد بامد	ا دا د	. L I				ıyl g														+
two-carbon diacid and a	i one-co	arbon d	liaico	noı.	_			l pro re ar				. •										on
two-carbon dialcohol an	d a ona	-carbo	n dia	ıcid				e a c					•									
	u u one	-cui bo	II GIO	icia.		_	_	the					0110		u.15				1440	' '',	ai ox	7
Propanal and ethanol.					9'	Jupo		,,,,	,g.	Ju		••										+
a chemist is analyzing a polymer is the chemist more an addition polymer. a condensation polymer. a polymer made from alker a polymer with carbon-car	Conder contair of a sr conder	y analy nsation ning tw mall mo nsation	poly poly o fun blecu poly	emer nction le, s mer.	s, su onal	ch a grou	s po	olyest leadi	ers,	are	for e cr	mec eati	l th	roug of es	h th	ne ro link	eact age	tion s ar	of nd t	mor he	nome relea	as
polyester is form	ed th	moug	h c	ביחם.	den	Sad	cior	,, 0	he	re	N	aul	er:	g	et:	5 0	70	du	ice	d	as	(
4 - 1																						_
Which of the following polymers is formed via addi vinyl chloride $(CH_2 = CHCl)$?	ition polymeriza	ation from the	monome	r										ers would						g unit -[
	ition polymeriza	ation from the	monome	r						CH_2 -										g unit -(-	
vinyl chloride ($CH_2=CHCl$)?	tion polymerize	ation from the	monome	r —						СН2 —	· CH(C									g unit -(
vinyl chloride ($CH_2=CHCl$)? $ \begin{tabular}{ c c c c c c c c c c c c c c c c c c c$	tion polymeriza	ation from the	monome	r						CH ₂ −	utane	$^{\circ}H_{2}CH$								g unit -(
vinyl chloride ($CH_2=CHCl$)?	ition polymerize	ation from the	monome							CH ₂ −	· CH(C	$^{\circ}H_{2}CH$								g unit -[
vinyl chloride ($CH_2=CHCl$)? $ \begin{tabular}{ c c c c c c c c c c c c c c c c c c c$	tion polymerize	ation from the	monome							CH ₂ −	utane	$^{\circ}H_{2}CH$								g unit -(
vinyl chloride ($CH_2=CHCl$)?	tion polymerize	ation from the	monome						Q	CH ₂ — Bi Bi C 2- Explanae	utane utene utene -Butene	e	[3)]- <i>n</i> -, ε		additio	n polyme	erizatio	n occur:	s?			

Chapter 20 Naming Carboxylic Acids Practice Worksheet

Carboxylic Acids:

Carboxylic Acids are named by counting the number of carbons in the logest continuous chain including the carboxyl group (—COOH) and by replacing the suffix '-ane' of the corresponding alkane with '-anoic acid.' If there are two —COOH groups, the suffix is expanded to include a prefix that indicates the number of —COOH groups present (-anedioic acid — there should not be more than 2 of these croups on the parent chain as they must occur at the ends). It is not necessary to indicate the position of the —COOH group because this group will be at the end of the parent chain and its carbon is automatically assigned as C-1.

1. Complete the exercise below by entering the formulas and condensed structural formulas of the first ten aldehydes.

		Carboxylic Acid Name	Condensed Structural Formula	
	a.	methanoic acid	СООН О	CHOZ
2	b.	Ethanoic acid	HC-C 0-H	C2H402
3	c.	proponois acid	CH3 CH2 = "101	1 C3H602
	d.	butanoic acid	CH ₃ -CH ₂ -CH ₂ -COOH	C44802
	e.	<u>Pentanoic</u> acid	CH3-CH2-CH2-CH2 -C	00H C5H10O2
	f.	Hexanoic acid		
	g.	Heptanoic acid		
	h.	Octanoic acid		
	i.	Nonanoic acid	CqH1802	
	j.	Decanoic acid	C ₁₀ H ₂₀ O ₂	

IUPAC Rules for Naming Carboxylic Acids:

RULE 1: The carboxyl group takes precedence over alkyl groups and halogen substituents, as well as double bonds, in the number of the parent chain.

$$\begin{array}{c|c} CH_{3} \\ \hline CH_{2} - CH_{2} - CH_{2} - CH_{2} \\ \hline CH_{2} - CH_{2} - CH_{2} - CH_{2} \\ \hline \end{array}$$

O
$$CH_2 - CH_3$$
HO $-C - CH - CH - CH_2 - CH_3$
 $-CH_2 - CH_3$

2,3-diethylhexanoic acid

4-ethyl-2-butylheptanoic acid

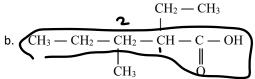
2. For the following compounds, draw a box around the longest continuous carbon chain and name each molecule. Number the carbons in the longest chain so that the alkyl group(s) will be on the <u>lowest</u> numbered carbons. The first one is done for you.

a.
$$CH_3 - CH_2 - CH - C - OH$$

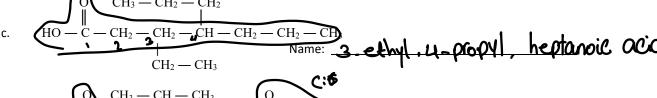
$$CH_3$$

$$CH_2 - CH$$

ame: 2-methylhut



Name: 1- ethyl, 2-methyl, pentonoic oc



d. HO C C CH₂ C C CH CH₂ C OH Name: 3-methy 3-propyl hexand CH₃

Name: _____

$$\begin{array}{c|c} CH_3 & CH_2-CH_2-CH_2-CH_3\\ & & \\ | & & \\ \text{f.} & CH_3-CH-CH-C-OH \\ & & \\ & & \\ O \end{array}$$

Name:

RULE 2: If the carboxyl group is attached to a ring, the parent ring is named and the suffix -carboxylic acid is added.

RULE 3: When both double bonds and carboxyl groups are present, the -en suffix follows the parent chain directly and the -oic acid suffix follows the -en suffix (notice that the e is left off, -en instead of -ene). The location of the double bond(s) is(are) indicated before the parent name as before, and the -oic acid suffix follows the -en suffix directly. Remember it is not necessary to specify the location of the carboxyl group because it will automatically be carbon #1. See below for examples. Again, the carboxyl gets priority in the numbering of the parent chain.

$$\begin{array}{c|ccccc} CH_3 & O & CH_2-CH_3 \\ & & & & \\ CH_3-CH_2=C-CH-C-OH & HO-C-C=CH-C-OH \\ & & & & \\ CH_3 & O & O \end{array}$$

2,3-dimethyl-3-pentenal

2-ethyl-2-butenedioic acid

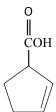
RULE 4: If there is a choice in numbering not previously covered, the parent chain is numbered to give the substituents the **lowest** number at the **first point of difference**.

$$CH_{3}$$
 $CH = CH - CH_{3}$ $CH_{3} - CH = C - C - OH$ $CH_{3} - CH - C - OH$ $CH_{3} - CH - C - OH$ $CH_{3} - CH - C - OH$

$$CH = CH - CH_{2}$$

$$CH_{3} - CH - C - OH$$

$$0$$



2-methyl-2-butenoic acid

2-methyl-3-pentenoic acid

2-cyclo-3-pentenoic acid

3. For the following compounds, draw a box around the longest continuous carbon chain and name each molecule. Number the carbons in the longest chain so that the alkyl group(s) will be on the lowest numbered carbons. The first one is done for you.

a.
$$CH_3 - CH = CH - C - CH - CH - CH_2 - C - OH$$

$$\begin{vmatrix} & & & \\ & &$$

$$\begin{array}{c} O & CH_2-CH_3 \\ \parallel & | \\ \text{d.} & HO-C-CH_2=CH_2-CH-CH_2-CH-CH_3} \\ | & \\ CH_2-CH_3 \\ O & \text{Name:} \end{array}$$

e.
$$CH_2 = CH - C - OH$$

Name:

Name:

$$O = COH$$



Name:

- 4. Draw the following compounds.
 - a. 4-methylhexanoic acid
 - b. 3-methylbutanoic acid
 - c. 2,6-dimethylheptanoic acid
 - d. 2-methyl-3-isopropylpentanoic acid
 - e. 3-ethyl-6-methyl-3-octenoic acid
 - f. 4-methyl-2-pentenoic acid
 - g. 3-ethyl-5-nonenoic acid
 - h. 3-ethyl-2-cyclopentanoic acid

Organic Chemistry - HYDROCARBONS WORKSHEET

Draw the complete structural formula and condensed molecular formula for each compound.

IUPAC Name	Complete Structural Formula	Condensed Formula
Single butane Caff AC		CAHIO
Single bond alkane: 2 methylhexane 2H3 C	4C-CH-CH-CH-CH-CH-3 CH3	C7 HIG
3-ethyl-2-methylnonane	CH3 CH3-CH3-CH3-CH3-CH3	C12 H26
alkere: double loand propene	$H \subset = CH - CH^3$	C3H6
4-methyl-2-heptene	H: 16 CH3- CH=CH-CH-CH2-CH2-CH3	Cg HIG
<u>etn</u> yne	H-C ≡ C-H	C2H2 n 2n-2
5,6-dimethyl-2-octyne	H-C-HH H-C-HH H-C-HH	C10 H18

cyclopropane 3	M2 CH2	C ₃ H ₆
1,3-diethylcyclohexane	HC O OCH - CH - CH S	C10 H20
5-ethyl-4-propyl-2- heptyne	HC -C≡C-CH-CH-CH2-CH3 CH2 CH2 CH3 CH3	C12 H 22
2,2-dimethly-3-phenylhexane	CH3 CH-CH-CH-CH-C	15 C14 H22
1-methyl-3- propylbenzene	CH CH CH 2 CH 2 CH 2 CH 3 CH 3 CH 3 CH 3	C10 H14

Draw structural formulae and give the names for the five possible (noncyclic) isomers of C_6H_{14} .