

Connections

Building on:

- This chapter does not depend on Chapter 1

Leading to:

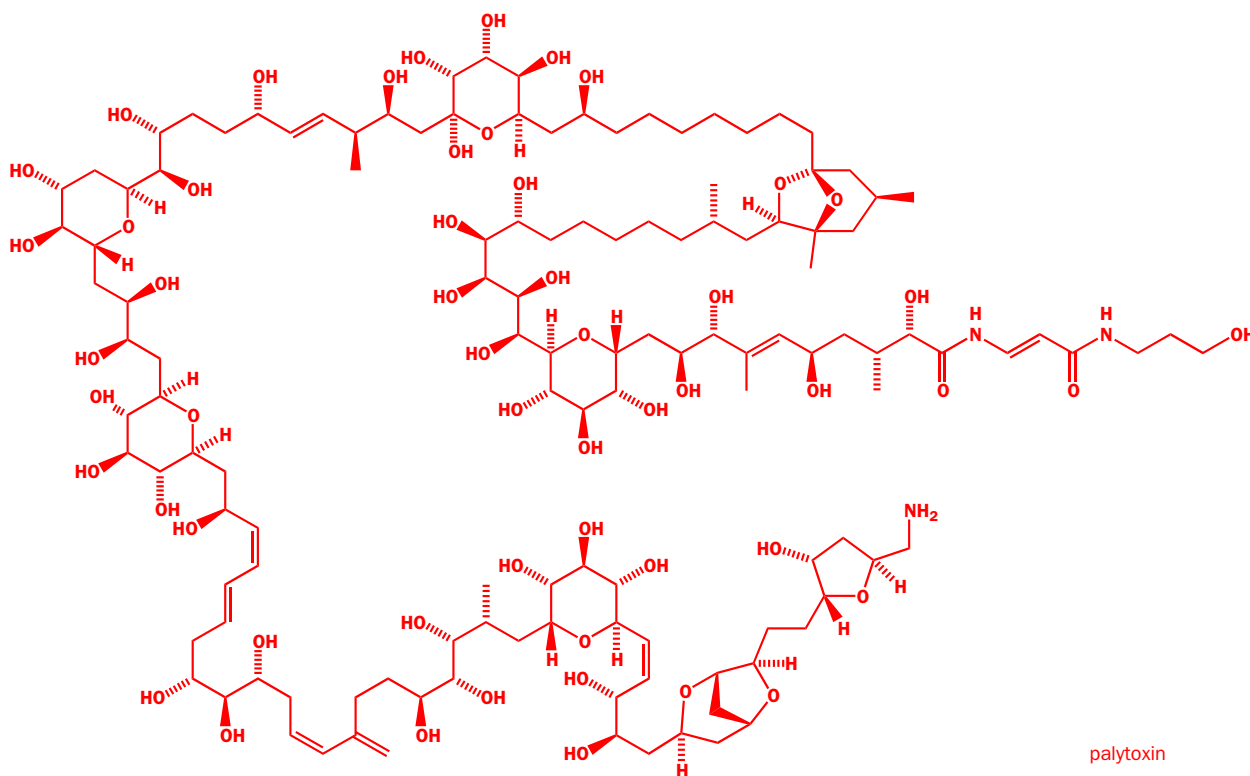
- The diagrams used in the rest of the book
- Why we use these particular diagrams
- How organic chemists name molecules in writing and in speech
- What is the skeleton of an organic molecule
- What is a functional group
- Some abbreviations used by all organic chemists
- Drawing organic molecules realistically in an easily understood style

Looking forward to:

- Ascertaining molecular structure spectroscopically **ch3**
- What determines a molecule's structure **ch4**

There are over 100 elements in the periodic table. Many molecules contain well over 100 atoms—palytoxin, for example (a naturally occurring compound with potential anticancer activity) contains 129 carbon atoms, 221 hydrogen atoms, 54 oxygen atoms, and 3 nitrogen atoms. It's easy to see how chemical structures can display enormous variety, providing enough molecules to build even the most complicated living creatures. But how can we understand what seems like a recipe for confusion? Faced with the collection of atoms we call a molecule, how can we make sense of what we see? This chapter will teach you how to interpret organic structures. It will also teach you how to draw organic molecules in a way that conveys all the necessary information and none of the superfluous.

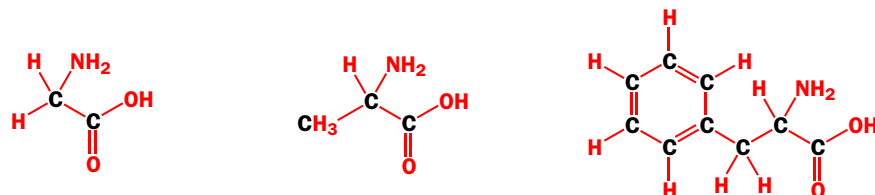
■ Palytoxin was isolated in 1971 in Hawaii from *Limu make o Hane* ('deadly seaweed of Hana') which had been used to poison spear points. It is one of the most toxic compounds known requiring only about 0.15 microgram per kilogram for death by injection. The complicated structure was determined a few years later.



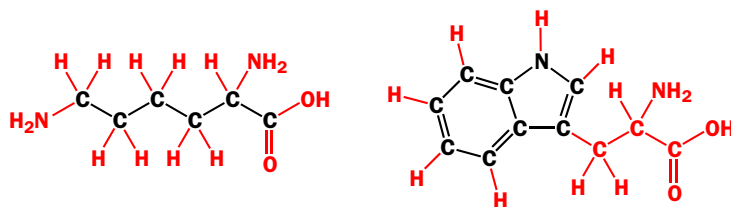
Hydrocarbon frameworks and functional groups

As we explained in Chapter 1, organic chemistry is the study of compounds that contain carbon. Nearly all organic compounds also contain hydrogen; most also contain oxygen, nitrogen, or other elements. Organic chemistry concerns itself with the way in which these atoms are bonded together into stable molecular structures, and the way in which these structures change in the course of chemical reactions.

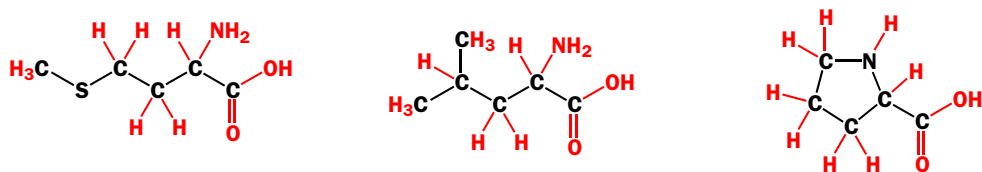
Some molecular structures are shown below. These molecules are all amino acids, the constituents of proteins. Look at the number of carbon atoms in each molecule and the way they are bonded together. Even within this small class of molecules there's great variety—*glycine* and *alanine* have only two or three carbon atoms; *phenylalanine* has nine.



Lysine has a chain of atoms; *tryptophan* has rings.



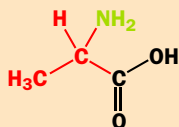
In *methionine* the atoms are arranged in a single chain; in *leucine* the chain is branched. In *proline*, the chain bends back on itself to form a ring.



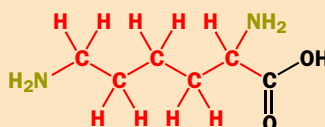
■ We shall return to amino acids as examples several times in this chapter, but we shall leave detailed discussions about their chemistry till Chapters 24 and 49, when we look at the way in which they polymerize to form peptides and proteins.

Yet all of these molecules have similar properties—they are all soluble in water, they are all both acidic and basic (amphoteric), they can all be joined with other amino acids to form proteins. This is because the chemistry of organic molecules depends much less on the number or the arrangement of carbon or hydrogen atoms than on the other types of atoms (O, N, S, P, Si...) in the molecule. We call parts of molecules containing small collections of these other atoms **functional groups**, simply because they are groups of atoms that determine the way the molecule works. All amino acids contain two functional groups: an amino (NH₂ or NH) group and a carboxylic acid (CO₂H) group (some contain other functional groups as well).

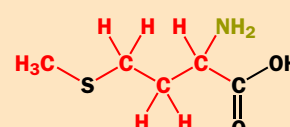
● The **functional groups** determine the way the molecule works both chemically and biologically.



alanine
contains just the amino
and carboxylic acid
functional groups



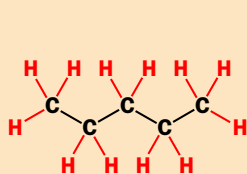
lysine
has an additional
amino group



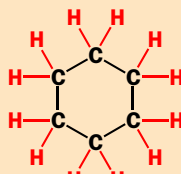
methionine
also has a sulfide
functional group

That isn't to say the carbon atoms aren't important; they just play quite a different role from those of the oxygen, nitrogen, and other atoms they are attached to. We can consider the chains and rings of carbon atoms we find in molecules as their skeletons, which support the functional groups and allow them to take part in chemical interactions, much as your skeleton supports your internal organs so they can interact with one another and work properly.

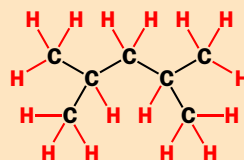
- The **hydrocarbon framework** is made up of chains and rings of carbon atoms, and it acts as a support for the functional groups.



a chain



a ring



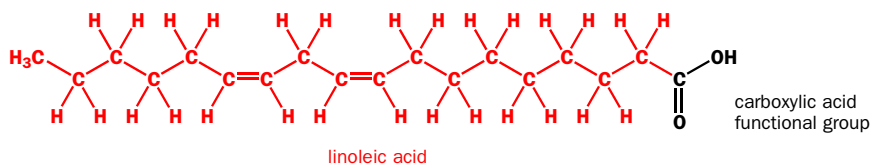
a branched chain

We will see later how the interpretation of organic structures as hydrocarbon frameworks supporting functional groups helps us to understand and rationalize the reactions of organic molecules. It also helps us to devise simple, clear ways of representing molecules on paper. You saw in Chapter 1 how we represented molecules on paper, and in the next section we shall teach you ways to draw (and ways not to draw) molecules—the handwriting of chemistry. *This section is extremely important*, because it will teach you how to communicate chemistry, clearly and simply, throughout your life as a chemist.

Drawing molecules

Be realistic

Below is another organic structure—again, you may be familiar with the molecule it represents; it is a fatty acid commonly called linoleic acid.

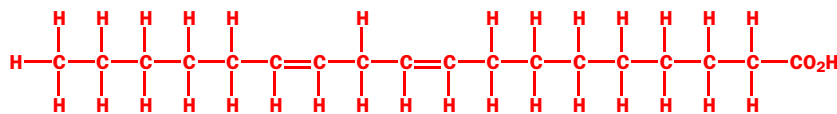


We could also depict linoleic acid as



linoleic acid

or as



linoleic acid

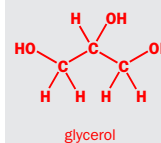
You may well have seen diagrams like these last two in older books—they used to be easy to print (in the days before computers) because all the atoms were in a line and all the angles were 90°. But are they realistic? We will consider ways of determining the shapes and structures of molecules in more detail in Chapter 3, but the picture below shows the structure of linoleic acid determined by X-ray crystallography.

Organic skeletons

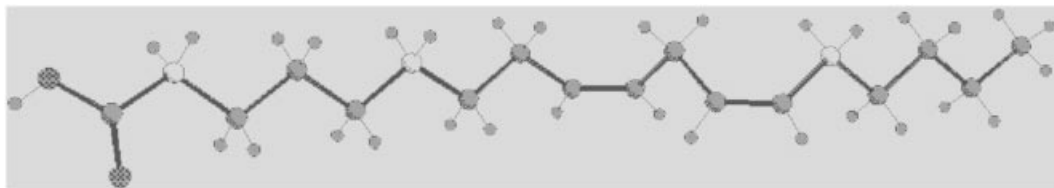
Organic molecules left to decompose for millions of years in the absence of light and oxygen become literally carbon skeletons—crude oil, for example, is a mixture of molecules consisting of nothing but carbon and hydrogen, while coal consists of little else but carbon. Although the molecules in coal and oil differ widely in chemical structure, they have one thing in common: no functional groups! Many are very unreactive: about the only chemical reaction they can take part in is combustion, which, in comparison to most reactions that take place in chemical laboratories or in living systems, is an extremely violent process. In Chapter 5 we will start to look at the way that functional groups direct the chemical reactions of a molecule.

Three fatty acid molecules and one glycerol molecule combine to form the fats that store energy in our bodies and are used to construct the membranes around our cells. This particular fatty acid, linoleic acid, cannot be manufactured in the human body, and is an essential part of a healthy diet found, for example, in sunflower oil.

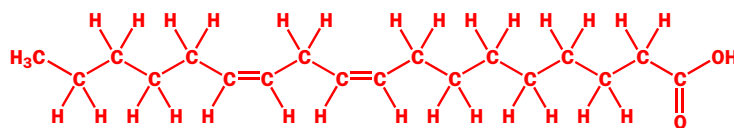
Fatty acids differ in the length of their chains of carbon atoms, yet they have very similar chemical properties because they all contain the carboxylic acid functional group. We shall come back to fatty acids in Chapter 49.



X-ray crystallography discovers the structures of molecules by observing the way X-rays bounce off atoms in crystalline solids. It gives clear diagrams with the atoms marked as circles and the bonds as rods joining them together.



You can see that the chain of carbon atoms is not linear, but a zig-zag. Although our diagram is just a two-dimensional representation of this three-dimensional structure, it seems reasonable to draw it as a zig-zag too.



linoleic acid

This gives us our first guideline for drawing organic structures.

● Guideline 1

Draw chains of atoms as zig-zags

Realism of course has its limits—the X-ray structure shows that the linoleic acid molecule is in fact slightly bent in the vicinity of the double bonds; we have taken the liberty of drawing it as a ‘straight zig-zag’. Similarly, close inspection of crystal structures like this reveals that the angle of the zig-zag is about 109° when the carbon atom is not part of a double bond and 120° when it is. The 109° angle is the ‘tetrahedral angle’, the angle between two vertices of a tetrahedron when viewed from its centre. In Chapter 4 we shall look at why carbon atoms take up this particular arrangement of bonds. Our realistic drawing is a projection of a three-dimensional structure onto flat paper so we have to compromise.

Be economical

When we draw organic structures we try to be as realistic as we can be without putting in superfluous detail. Look at these three pictures.



1



2

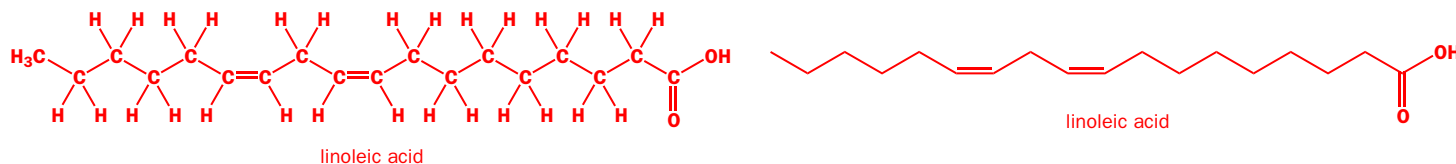


3

(1) is immediately recognizable as Leonardo da Vinci’s Mona Lisa. You may not recognize (2)—it’s also Leonardo da Vinci’s Mona Lisa—this time viewed from above. The frame is very ornate, but the picture tells us as much about the painting as our rejected linear and 90° angle diagrams did about

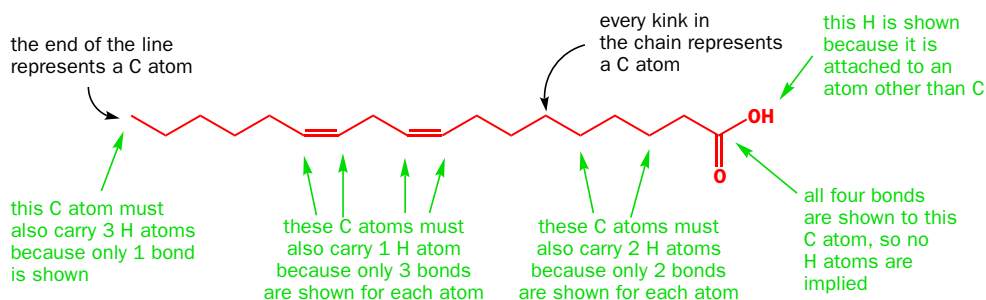
our fatty acid. They're both correct—in their way—but sadly useless. What we need when we draw molecules is the equivalent of (3). It gets across the idea of the original, and includes all the detail necessary for us to recognize what it's a picture of, and leaves out the rest. And it was quick to draw—this picture was drawn in less than 10 minutes: we haven't got time to produce great works of art!

Because functional groups are the key to the chemistry of molecules, clear diagrams must emphasize the functional groups, and let the hydrocarbon framework fade into the background. Compare the diagrams below:



The second structure is the way that most organic chemists would draw linoleic acid. Notice how the important carboxylic acid functional group stands out clearly and is no longer cluttered by all those Cs and Hs. The zig-zag pattern of the chain is much clearer too. And this structure is much quicker to draw than any of the previous ones!

To get this diagram from the one above we've done two things. Firstly, we've got rid of all the hydrogen atoms attached to carbon atoms, along with the bonds joining them to the carbon atoms. Even without drawing the hydrogen atoms we know they're there—we assume that any carbon atom that doesn't appear to have its potential for four bonds satisfied is also attached to the appropriate number of hydrogen atoms. Secondly, we've rubbed out all the Cs representing carbon atoms. We're left with a zig-zag line, and we assume that every kink in the line represents a carbon atom, as does the end of the line.



We can turn these two simplifications into two more guidelines for drawing organic structures.

● Guideline 2

Miss out the Hs attached to carbon atoms, along with the C–H bonds (unless there is a good reason not to)

● Guideline 3

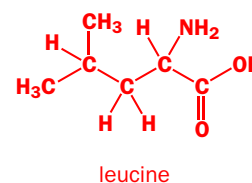
Miss out the capital Cs representing carbon atoms (unless there is a good reason not to)

▶ What is 'a good reason not to'? One is if the C or H is part of a functional group. Another is if the C or H needs to be highlighted in some way, for example, because it's taking part in a reaction. Don't be too rigid about these guidelines: they're not rules. Better is just to learn by example (you'll find plenty in this book): if it helps clarify, put it in; if it clutters and confuses, leave it out. One thing you must remember, though: if you write a carbon atom as a letter C then you *must* add all the H atoms too. If you don't want to draw all the Hs, don't write C for carbon.

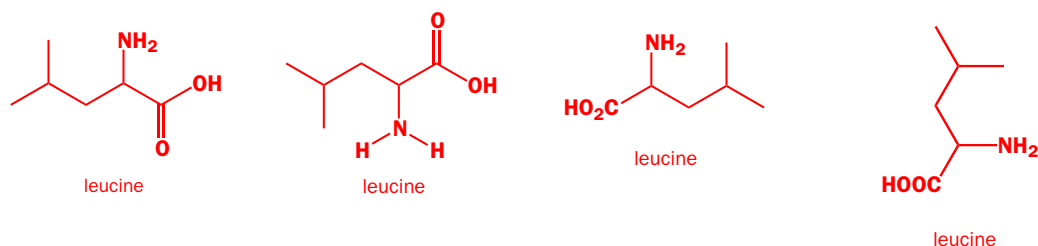
Be clear

Try drawing some of the amino acids represented on p. 000 in a similar way, using the three guidelines. The bond angles at tetrahedral carbon atoms are about 109° . Make them look about 109° projected on to a plane! (120° is a good compromise, and it makes the drawings look neat.)

Start with leucine — earlier we drew it as the structure to the right. Get a piece of paper and do it now; then see how your drawing compares with our suggestions.

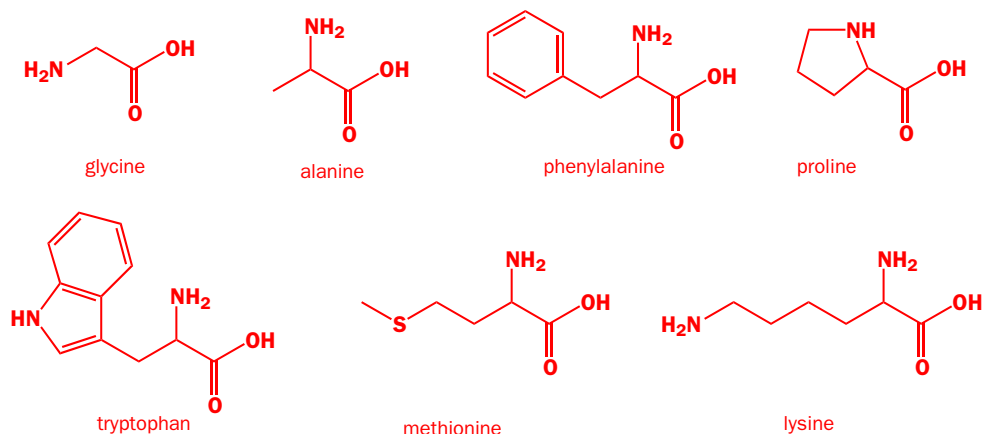


It doesn't matter which way up you've drawn it, but your diagram should look something like one of these structures below.



The guidelines we gave were only guidelines, not rules, and it certainly does not matter which way round you draw the molecule. The aim is to keep the functional groups clear, and let the skeleton fade into the background. That's why the last two structures are all right—the carbon atom shown as 'C' is part of a functional group (the carboxyl group) so it can stand out.

Now turn back to p. 000 and try redrawing the some of the other eight structures there using the guidelines. Don't look at our suggestions below until you've done them! Then compare your drawings with our suggestions.

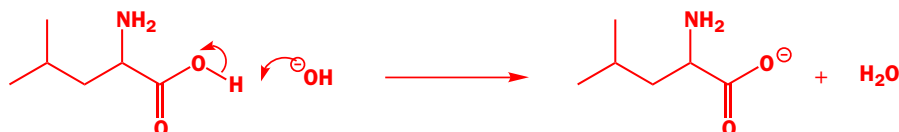


Remember that these are only suggestions, but we hope you'll agree that this style of diagram looks much less cluttered and makes the functional groups much clearer than the diagrams on p. 000. Moreover, they still bear significant resemblance to the 'real thing'—compare these crystal structures of lysine and tryptophan with the structures shown above, for example.

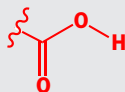
Structural diagrams can be modified to suit the occasion

You'll probably find that you want to draw the same molecule in different ways on different occasions to emphasize different points. Let's carry on using leucine as an example. We mentioned before that an amino acid can act as an acid or as a base. When it acts as an acid, a base (for example, hydroxide, OH^-) removes H^+ from the carboxylic acid group in a reaction we can represent as

■ Not all chemists put circles round their plus and minus charges—it's a matter of personal choice.



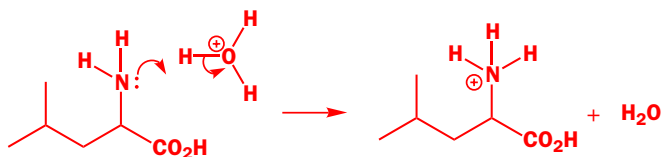
■ The wiggly line is a graphical way of indicating an incomplete structure: it shows where we have mentally 'snapped off' the CO_2H group from the rest of the molecule.



The product of this reaction has a negative charge on an oxygen atom. We have put it in a circle to make it clearer, and we suggest you do the same when you draw charges: $+$'s and $-$'s are easily mislaid. We shall discuss this type of reaction, the way in which reactions are drawn, and what the 'curly arrows' in the diagram mean in Chapter 5. But for now, notice that we drew out the CO_2H as the fragment left because we wanted to show how the $\text{O}-\text{H}$ bond was broken when the base attacked. We modified our diagram to suit our own purposes.

When leucine acts as a base, the amino (NH_2) group is involved. The nitrogen atom attaches itself to a proton, forming a new bond using its *lone pair*.

We can represent this reaction as



Notice how we drew the lone pair at this time because we wanted to show how it was involved in the reaction. The oxygen atoms of the carboxylic acid groups also have lone pairs but we didn't draw them in because they weren't relevant to what we were talking about. Neither did we feel it was necessary to draw CO_2H in full this time because none of the atoms or bonds in the carboxylic acid functional group was involved in the reaction.

Structural diagrams can show three-dimensional information on a two-dimensional page

Of course, all the structures we have been drawing only give an idea of the real structure of the molecules. For example, the carbon atom between the NH_2 group and the CO_2H group of leucine has a tetrahedral arrangement of atoms around it, a fact which we have so far completely ignored.

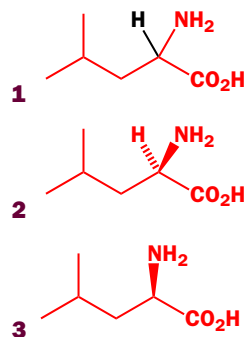
We might want to emphasize this fact by drawing in the hydrogen atom we missed out at this point as in structure 1 (in the right-hand margin).

We can then show that one of the groups attached to this carbon atom comes towards us, out of the plane of the paper, and the other one goes away from us, into the paper. There are several ways of doing this. In structure 2, the bold, wedged bond suggests a perspective view of a bond coming towards you, while the hashed bond suggests a bond fading away from you. The other two 'normal' bonds are in the plane of the paper.

Alternatively we could miss out the hydrogen atom and draw something a bit neater though slightly less realistic as structure 3.

We can assume the missing hydrogen atom is behind the plane of the paper, because that is where the 'missing' vertex of the tetrahedron of atoms attached to the carbon atom lies. These conventions allow us to give an idea of the three-dimensional shape (stereochemistry) of any organic molecule—you have already seen them in use in the diagram of the structure of palytoxin at the beginning of this chapter.

■ A lone pair is a pair of electrons that is not involved in a chemical bond. We shall discuss lone pairs in detail in Chapter 4. Again, don't worry about what the curly arrows in this diagram mean—we will cover them in detail in Chapter 5.



▶ When you draw diagrams like these to indicate the three-dimensional shape of the molecule, try to keep the hydrocarbon framework in the plane of the paper and allow functional groups and other branches to project forwards out of the paper or backwards into it.

■ We shall look in more detail at the shapes of molecules—their *stereochemistry*—in Chapter 16.

● Reminder

Organic structures should be drawn to be *realistic, economical, clear*.

We gave three guidelines to help you achieve this when you draw structures:

- Guideline 1: Draw chains of atoms as *zig-zags*
- Guideline 2: Miss out the Hs attached to carbon atoms, along with the C–H bonds
- Guideline 3: Miss out the capital Cs representing carbon atoms

The guidelines we have given and conventions we have illustrated in this section have grown up over decades. They are used by organic chemists because they work! We guarantee to follow them for the rest of the book—try to follow them yourself whenever you draw an organic structure. Before you ever draw a capital C or a capital H again, ask yourself whether it's really necessary!

Now that we have considered how to draw structures, we can return to some of the structural types that we find in organic molecules. Firstly, we'll talk about hydrocarbon frameworks, then about functional groups.

Hydrocarbon frameworks

Carbon as an element is unique in the variety of structures it can form. It is unusual because it forms strong, stable bonds to the majority of elements in the periodic table, including itself. It is this ability to form bonds to itself that leads to the variety of organic structures that exist, and indeed to the possibility of life existing at all. Carbon may make up only 0.2% of the earth's crust, but it certainly deserves a whole branch of chemistry all to itself.

Chains

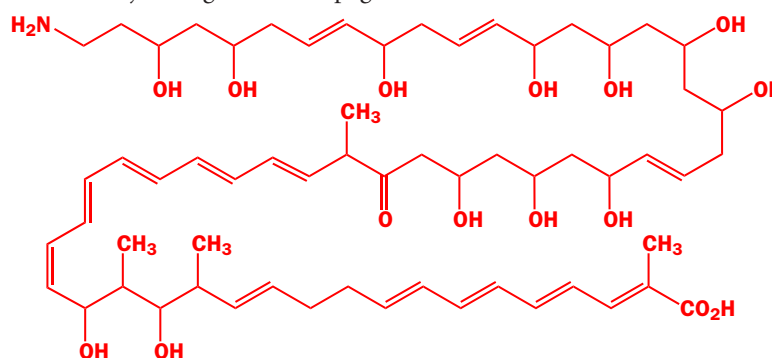
The simplest class of hydrocarbon frameworks contains just chains of atoms. The fatty acids we met earlier have hydrocarbon frameworks made of zig-zag chains of atoms, for example. Polythene is a polymer whose hydrocarbon framework consists entirely of chains of carbon atoms.



a section of the structure of polythene

At the other end of the spectrum of complexity is this antibiotic, extracted from a fungus in 1995 and aptly named linearmycin as it has a long linear chain. The chain of this antibiotic is so long that we have to wrap it round two corners just to get it on the page.

We haven't drawn whether the CH_3 groups and OH groups are in front of or behind the plane of the paper, because (at the time of writing this book) no one yet knows. The stereochemistry of linearmycin is unknown.



▶ Notice we've drawn in four groups as CH_3 —we did this because we didn't want them to get overlooked in such a large structure. They are the only tiny branches off this long winding trunk.

Names for carbon chains

It is often convenient to refer to a chain of carbon atoms by a name indicating its length. You have probably met some of these names before in the names of the simplest organic molecules, the alkanes. There are also commonly used abbreviations for these names: these can be very useful in both writing about chemistry and in drawing chemical structures, as we shall see shortly.

Names and abbreviations for carbon chains

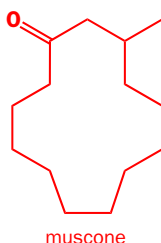
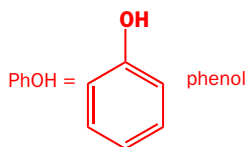
Number of carbon atoms in chain	Name of group	Formula [†]	Abbreviation	Name of alkane (= chain + H)
1	methyl	$-\text{CH}_3$	Me	methane
2	ethyl	$-\text{CH}_2\text{CH}_3$	Et	ethane
3	propyl	$-\text{CH}_2\text{CH}_2\text{CH}_3$	Pr	propane
4	butyl	$-(\text{CH}_2)_3\text{CH}_3$	Bu	butane
5	pentyl	$-(\text{CH}_2)_4\text{CH}_3$	— [‡]	pentane
6	hexyl	$-(\text{CH}_2)_5\text{CH}_3$	— [‡]	hexane
7	heptyl	$-(\text{CH}_2)_6\text{CH}_3$	— [‡]	heptane
8	octyl	$-(\text{CH}_2)_7\text{CH}_3$	— [‡]	octane
9	nonyl	$-(\text{CH}_2)_8\text{CH}_3$	— [‡]	nonane
10	decyl	$-(\text{CH}_2)_9\text{CH}_3$	— [‡]	decane

[†] This representation is not recommended.

[‡] Names for longer chains are not commonly abbreviated.

▶ The names for shorter chains (which you must learn) exist for historical reasons; for chains of 5 or more carbon atoms, the systematic names are based on Greek number names.

Of course, Ar = argon too, but so few argon compounds exist that there is never any confusion.

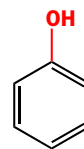


Any compound containing a benzene ring, or a related (Chapter 7) ring system is known as 'aromatic', and another useful organic element symbol related to Ph is Ar (for 'aryl'). While Ph always means C_6H_5 , Ar can mean any *substituted* phenyl ring, in other words, phenyl with any number of the hydrogen atoms replaced by other groups.

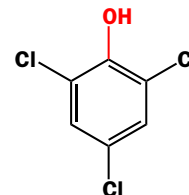
For example, while PhOH always means phenol, ArOH could mean phenol, 2,4,6-trichlorophenol (the antiseptic TCP), paracetamol or aspirin (among many other substituted phenols). Like R, the 'wild card' alkyl group, Ar is a 'wild card' *aryl* group.

The compound known as muscone has only relatively recently been made in the lab. It is the pungent aroma that makes up the base-note of musk fragrances. Before chemists had determined its structure and devised a laboratory synthesis the only source of musk was the musk deer, now rare for this very reason. Muscone's skeleton is a 13-membered ring of carbon atoms.

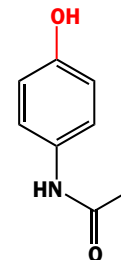
The steroid hormones have several (usually four) rings fused together. These are testosterone and oestradiol, the important human male and female sex hormones.



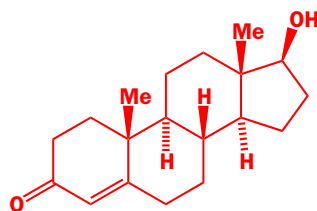
phenol



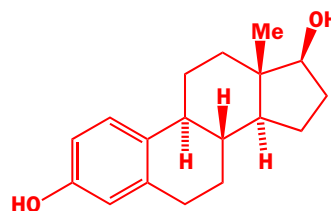
2,4,6-trichlorophenol



paracetamol

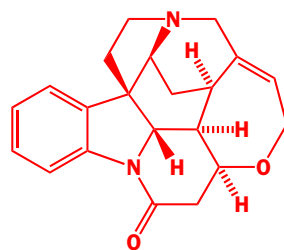


testosterone

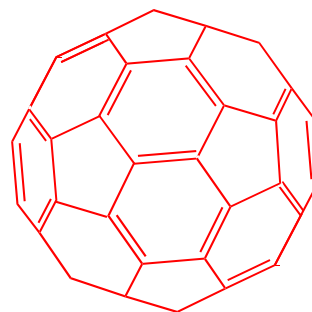


oestradiol

Some ring structures are much more complicated. The potent poison strychnine is a tangle of interconnecting rings.



strychnine



Buckminsterfullerene

Buckminsterfullerene

Buckminsterfullerene is named after the American inventor and architect Richard Buckminster Fuller, who designed the structures known as 'geodesic domes'.



One of the most elegant ring structures is shown above and is known as Buckminsterfullerene. It consists solely of 60 carbon atoms in rings that curve back on themselves to form a football-shaped cage.

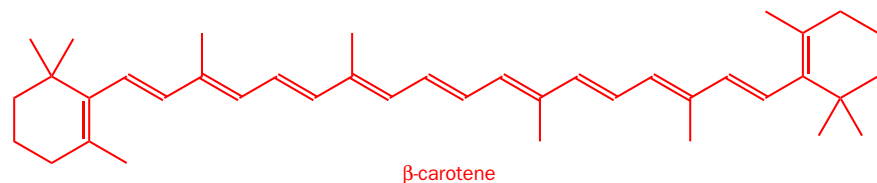
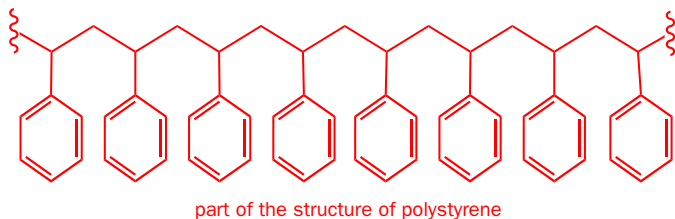
Count the number of bonds at any junction and you will see they add up to four so no hydrogens need be added. This compound is C_{60} . Note that you can't see all the atoms as some are behind the sphere.

Rings of carbon atoms are given names starting with 'cyclo', followed by the name for the carbon chain with the same number of carbon atoms.

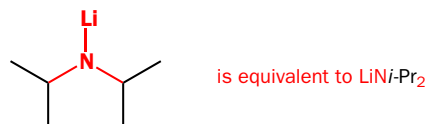
To the right, structure 1 shows chrysanthemic acid, part of the naturally occurring pesticides called pyrethrins (an example appears in Chapter 1), which contains a cyclopropane ring. Propane has three carbon atoms. Cyclopropane is a three-membered ring. Grandisol (structure 2), an insect pheromone used by male boll weevils to attract females, has a structure based on a cyclobutane ring. Butane has four carbon atoms. Cyclobutane is a four-membered ring. Cyclamate (structure 3), formerly used as an artificial sweetener, contains a cyclohexane ring. Hexane has six carbon atoms. Cyclohexane is a six-membered ring.

Branches

Hydrocarbon frameworks rarely consist of single rings or chains, but are often branched. Rings, chains, and branches are all combined in structures like that of the marine toxin palytoxin that we met at the beginning of the chapter, polystyrene, a polymer made of six-membered rings dangling from linear carbon chains, or of β -carotene, the compound that makes carrots orange.



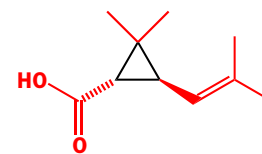
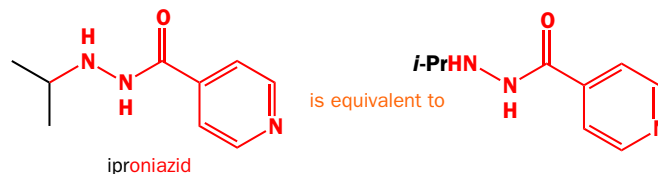
Just like some short straight carbon chains, some short branched carbon chains are given names and organic element symbols. The most common is the isopropyl group. Lithium diisopropylamide (also called LDA) is a strong base commonly used in organic synthesis.



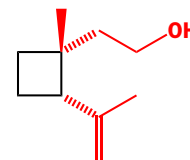
lithium diisopropylamide (LDA)

Iproniazid is an antidepressant drug with *i*-Pr in both structure and name.

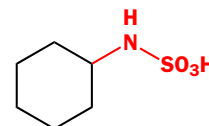
Notice how the 'propyl' part of 'isopropyl' still indicates three carbon atoms; they are just joined together in a different way—in other words, as an *isomer* of the straight chain propyl group. Sometimes, to avoid confusion, the straight chain alkyl groups are called '*n*-alkyl' (for example, *n*-Pr, *n*-Bu)—*n* for 'normal'—to distinguish them from their branched counterparts.



1 chrysanthemic acid



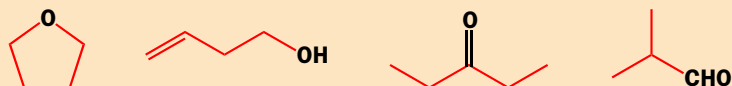
2 grandisol



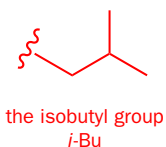
3 cyclamate



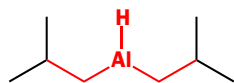
- **Isomers** are molecules with the same kinds and numbers of atoms joined up in different ways *n*-propanol, *n*-PrOH, and isopropanol, *i*-PrOH, are isomeric alcohols. Isomers need not have the same functional groups, these compounds are all isomers of $\text{C}_4\text{H}_8\text{O}$.



■ 'Isopropyl' may be abbreviated to *i*Pr, *i*Pr, or Pr^{*i*}. We will use the first in this book, but you may see the others used elsewhere.

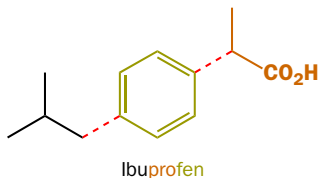


The isobutyl (*i*-Bu) group is a CH₂ group joined to an *i*-Pr group. It is *i*-PrCH₂–
Two isobutyl groups are present in the reducing agent diisobutyl aluminium hydride (DIBAL).

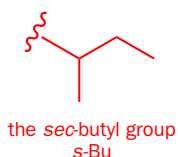


diisobutyl aluminium hydride (DIBAL)
is equivalent to **HAl*i*-Bu₂**

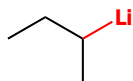
The painkiller ibuprofen (marketed as Nurofen[®]) contains an isobutyl group.



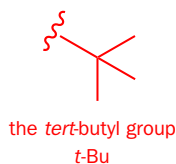
■ Notice how the invented name ibuprofen is a medley of 'ibu' (from *i*-Bu for isobutyl) + 'pro' (for propyl, the three-carbon unit shown in gold) + 'fen' (for the phenyl ring). We will talk about the way in which compounds are named later in this chapter.



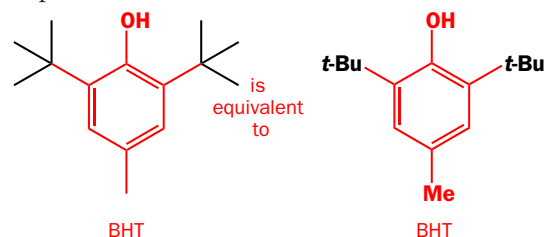
There are two more isomers of the butyl group, both of which have common names and abbreviations. The *sec*-butyl group (*s*-butyl or *s*-Bu) has a methyl and an ethyl group joined to the same carbon atom. It appears in an organolithium compound, *sec*-butyl lithium, used to introduce lithium atoms into organic molecules.



is equivalent to ***s*-BuLi**

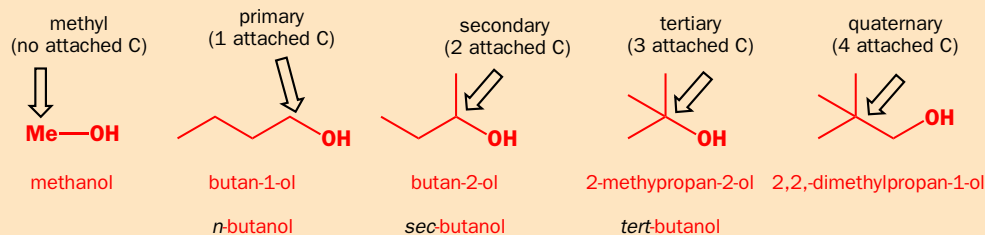


The *tert*-butyl group (*t*-butyl or *t*-Bu) group has three methyl groups joined to the same carbon atom. Two *t*-Bu groups are found in BHT ('butylated hydroxy toluene'), an antioxidant added to some processed foods.



● Primary, secondary, and tertiary

The prefixes *sec* and *tert* are really short for secondary and tertiary, terms that refer to the carbon atom that attaches these groups to the rest of the molecular structure.





A primary carbon atom is attached to only one other C atom, a secondary to two other C atoms, and so on. This means there are five types of carbon atom.

These names for bits of hydrocarbon framework are more than just useful ways of writing or talking about chemistry. They tell us something fundamental about the molecule and we shall use them when we describe reactions.

This quick architectural tour of some of the molecular edifices built by nature and by man serves just as an introduction to some of the hydrocarbon frameworks you will meet in the rest of this chapter and of this book. Yet, fortunately for us, however complicated the hydrocarbon framework might be, it serves only as a support for the functional groups. And, by and large, a functional group in one molecule behaves in much the same way as it does in another molecule. What we now need to do, and we start in the next section, is to introduce you to some functional groups, and to explain why it is that their attributes are the key to understanding organic chemistry.

Functional groups

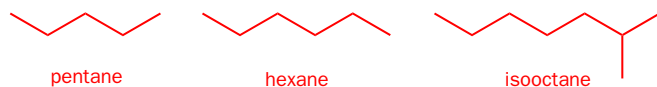
If you can take ethane gas (CH_3CH_3 , or EtH, or even , though a single line like this doesn't look much like a chemical structure) and bubble it through acids, bases, oxidizing agents, reducing agents—in fact almost any chemical you can think of—it will remain unchanged. Just about the only thing you can do with it is burn it. Yet ethanol ($\text{CH}_3\text{CH}_2\text{OH}$, or , or preferably EtOH) not only burns, it reacts with acids, bases, and oxidizing agents.

The difference between ethanol and ethane is the functional group—the OH or hydroxyl group. We know that these chemical properties (being able to react with acids, bases, and oxidizing agents) are properties of the hydroxyl group and not just of ethanol because other compounds containing OH groups (in other words, other alcohols) have similar properties, whatever their hydrocarbon frameworks.

Your understanding of functional groups will be the key to your understanding of organic chemistry. We shall therefore now go on to meet some of the most important functional groups. We won't say much about the properties of each group; that will come in Chapter 5 and later. Your task at this stage is to learn to recognize them when they appear in structures, so make sure you learn their names. The classes of compound associated with some functional groups also have names: for example, compounds containing the hydroxyl group are known as alcohols. Learn these names too as they are more important than the systematic names of individual compounds. We've told you a few snippets of information about each group to help you to get to know something of the group's character.

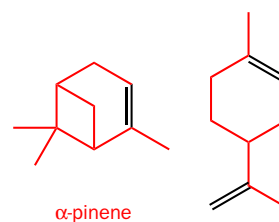
Alkanes contain no functional groups

The alkanes are the simplest class of organic molecules because they contain no functional groups. They are extremely unreactive, and therefore rather boring as far as the organic chemist is concerned. However, their unreactivity can be a bonus, and alkanes such as pentane and hexane are often used as solvents, especially for purification of organic compounds. Just about the only thing alkanes will do is burn—methane, propane, and butane are all used as domestic fuels, and petrol is a mixture of alkanes containing largely isooctane.



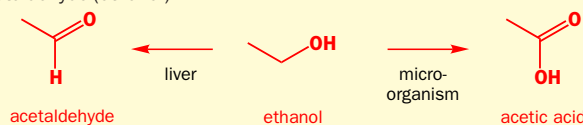
Alkenes (sometimes called olefins) contain C=C double bonds

It may seem strange to classify a type of bond as a functional group, but you will see later that C=C double bonds impart reactivity to an organic molecule just as functional groups consisting of, say, oxygen or nitrogen atoms do. Some of the compounds produced by plants and used by perfumers are alkenes (see Chapter 1). For example, pinene has a smell evocative of pine forests, while limonene smells of citrus fruits.



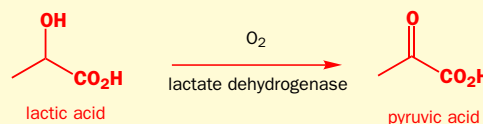
Ethanol

The reaction of ethanol with oxidizing agents makes vinegar from wine and sober people from drunk ones. In both cases, the oxidizing agent is oxygen from the air, catalysed by an enzyme in a living system. The oxidation of ethanol by microorganisms that grow in wine left open to the air leads to acetic acid (ethanoic acid) while the oxidation of ethanol by the liver gives acetaldehyde (ethanal).

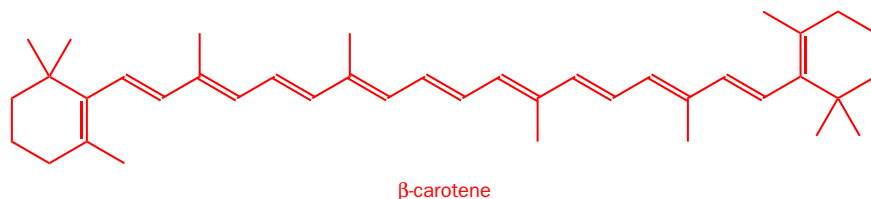


Human metabolism and oxidation

The human metabolism makes use of the oxidation of alcohols to render harmless other toxic compounds containing the OH group. For example, lactic acid, produced in muscles during intense activity, is oxidized by an enzyme called lactate dehydrogenase to the metabolically useful compound pyruvic acid.



You've already met the orange pigment β -carotene. Eleven C=C double bonds make up most of its structure. Coloured organic compounds often contain chains of C=C double bonds like this. In Chapter 7 you will find out why this is so.

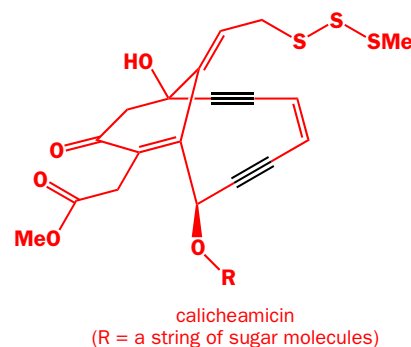


► Saturated and unsaturated carbon atoms

In an alkane, each carbon atom is joined to four other atoms (C or H). It has no potential for forming more bonds and is therefore **saturated**. In alkenes, the carbon atoms making up the C=C double bond are attached to only three atoms each. They still have the potential to bond with one more atom, and are therefore **unsaturated**. In general, carbon atoms attached to four other atoms are saturated; those attached to three, two, or one are unsaturated.

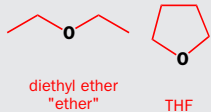
Alkynes contain C \equiv C triple bonds

Just like C=C double bonds, C \equiv C triple bonds have a special type of reactivity associated with them, so it's useful to call a C \equiv C triple bond a functional group. Alkynes are **linear** so we draw them with four carbon atoms in a straight line. Alkynes are not as widespread in nature as alkenes, but one fascinating class of compounds containing C \equiv C triple bonds is a group of antitumour agents discovered during the 1980s. Calicheamicin is a member of this group. The high reactivity of this combination of functional groups enables calicheamicin to attack DNA and prevent cancer cells from proliferating. For the first time we have drawn a molecule in three dimensions, with two bonds crossing one another—can you see the shape?



Remember that R can mean any alkyl group.

If we want a structure to contain more than one 'R', we give the R's numbers and call them R¹, R²... Thus R¹-O-R² means an ether with two different unspecified alkyl groups. (Not R₁, R₂..., which would mean 1 \times R, 2 \times R...)

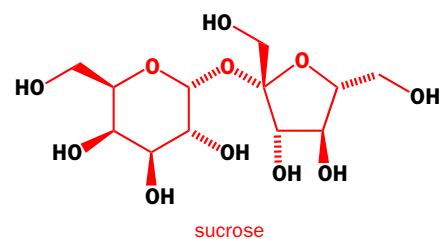


Another common laboratory solvent is called 'petroleum ether'. Don't confuse this with diethyl ether! Petroleum ether is in fact not an ether, but a mixture of alkanes. 'Ether', according to the *Oxford English Dictionary*, means 'clear sky, upper region beyond the clouds', and hence used to be used for anything light, airy, and volatile.

Alcohols (R-OH) contain a hydroxyl (OH) group

We've already talked about the hydroxyl group in ethanol and other alcohols. Carbohydrates are peppered with hydroxyl groups; sucrose has eight of them for example (a more three-dimensional picture of the sucrose molecule appears in Chapter 1).

Molecules containing hydroxyl groups are often soluble in water, and living things often attach sugar groups, containing hydroxyl groups, to otherwise insoluble organic compounds to keep them in solution in the cell. Calicheamicin, a molecule we have just mentioned, contains a string of sugars for just this reason. The liver carries out its task of detoxifying unwanted organic compounds by repeatedly hydroxylating them until they are water-soluble, and they are then excreted in the bile or urine.



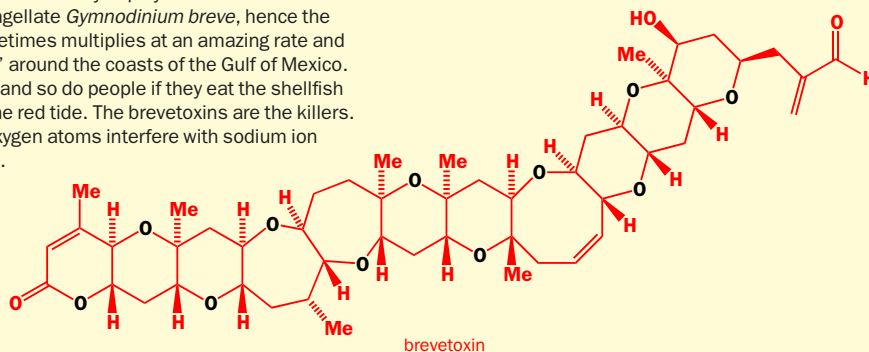
Ethers (R¹-O-R²) contain an alkoxy group (-OR)

The name **ether** refers to any compound that has two alkyl groups linked through an oxygen atom. 'Ether' is also used as an everyday name for diethyl ether, Et₂O. You might compare this use of the word 'ether' with the common use of the word 'alcohol' to mean ethanol. Diethyl ether is a highly flammable solvent that boils at only 35°C. It used to be used as an anaesthetic. Tetrahydrofuran (THF) is another commonly used solvent and is a cyclic ether.

Brevetoxin B is a fascinating naturally occurring compound that was synthesized in the laboratory in 1995. It is packed with ether functional groups in ring sizes from 6 to 8.

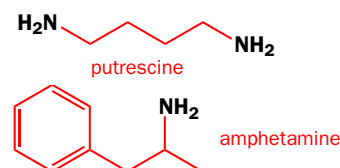
Brevetoxin B

Brevetoxin B is one of a family of polyethers found in a sea creature (a dinoflagellate *Gymnodinium breve*, hence the name) which sometimes multiplies at an amazing rate and creates 'red tides' around the coasts of the Gulf of Mexico. Fish die in shoals and so do people if they eat the shellfish that have eaten the red tide. The brevetoxins are the killers. The many ether oxygen atoms interfere with sodium ion (Na^+) metabolism.



Amines (R-NH_2) contain the amino (NH_2) group

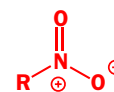
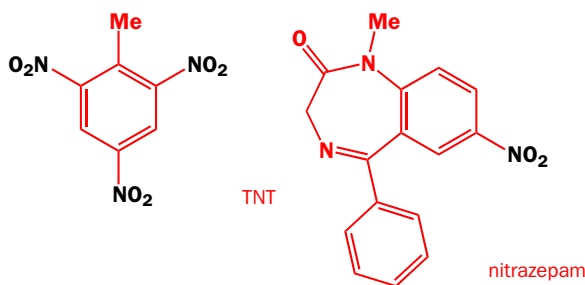
We met the amino group when we were discussing the amino acids: we mentioned that it was this group that gave these compounds their basic properties. Amines often have powerful fishy smells: the smell of putrescine is particularly foul. It is formed as meat decays. Many neurologically active compounds are also amines: amphetamine is a notorious stimulant.



Nitro compounds (R-NO_2) contain the nitro group (NO_2)

The nitro group (NO_2) is often incorrectly drawn with five bonds to nitrogen which you will see in Chapter 4, is impossible. Make sure you draw it correctly when you need to draw it out in detail. If you write just NO_2 you are all right!

Several nitro groups in one molecule can make it quite unstable and even explosive. Three nitro groups give the most famous explosive of all, TNT (trinitrotoluene), its kick.



the nitro group

nitrogen cannot have five bonds!

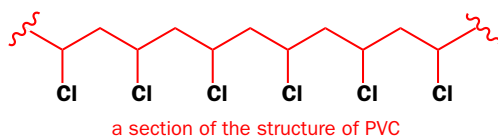


incorrect structure for the nitro group

However, functional groups refuse to be stereotyped. Nitrazepam also contains a nitro group, but this compound is marketed as Mogadon[®], the sleeping pill.

Alkyl halides (fluorides R-F , chlorides R-Cl , bromides R-Br , or iodides R-I) contain the fluoro, chloro, bromo, or iodo groups

These three functional groups have similar properties—though alkyl iodides are the most reactive and alkyl fluorides the least. PVC (polyvinyl chloride) is one of the most widely used polymers—it has a chloro group on every other carbon atom along a linear hydrocarbon framework. Methyl iodide (MeI), on the other hand, is a dangerous carcinogen, since it reacts with DNA and can cause mutations in the genetic code.



These compounds are also known as haloalkanes (fluoroalkanes, chloroalkanes, bromoalkanes or iodoalkanes).

Because alkyl halides have similar properties, chemists use yet another 'wild card' organic element, X, as a convenient substitute for Cl, Br, or I (sometimes F). So R-X is any alkyl halide.

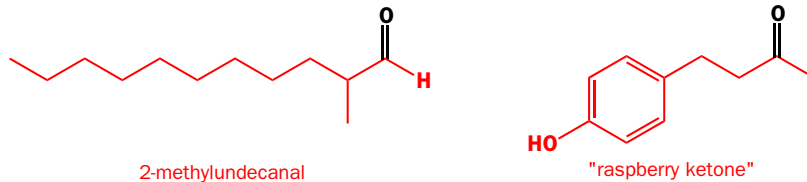
■
-CHO represents:



When we write aldehydes as R-CHO, we have no choice but to write in the C and H (because they're part of the functional group)—one important instance where you should ignore Guideline 3 for drawing structures. Another point: always write R-CHO and never R-COH, which looks too much like an alcohol.

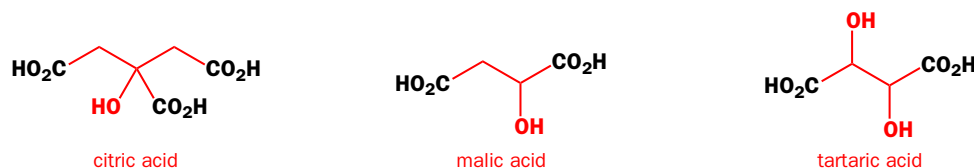
Aldehydes (R-CHO) and ketones (R¹-CO-R²) contain the carbonyl group C=O

Aldehydes can be formed by oxidizing alcohols—in fact the liver detoxifies ethanol in the bloodstream by oxidizing it first to acetaldehyde (ethanal, CH₃CHO). Acetaldehyde in the blood is the cause of hang-overs. Aldehydes often have pleasant smells—2-methylundecanal is a key component of the fragrance of Chanel No 5™, and 'raspberry ketone' is the major component of the flavour and smell of raspberries.



Carboxylic acids (R-CO₂H) contain the carboxyl group CO₂H

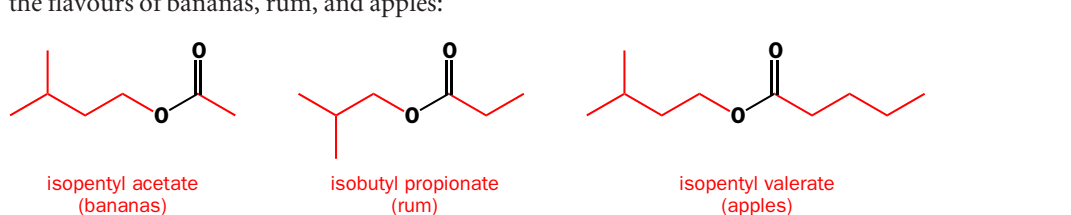
As their name implies, compounds containing the carboxylic acid (CO₂H) group can react with bases, losing a proton to form carboxylate salts. Edible carboxylic acids have sharp flavours and several are found in fruits—citric, malic, and tartaric acids are found in lemons, apples, and grapes, respectively.



Esters (R¹-CO₂R²) contain a carboxyl group with an extra alkyl group (CO₂R)

Fats are esters; in fact they contain three ester groups. They are formed in the body by condensing glycerol, a compound with three hydroxyl groups, with three fatty acid molecules.

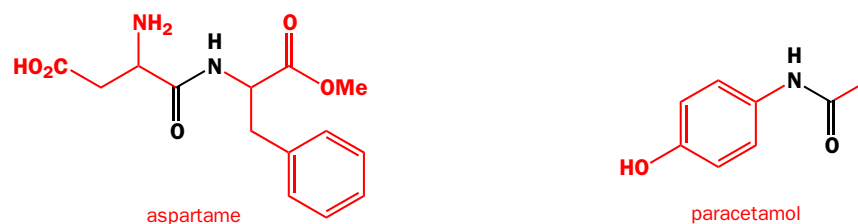
Other, more volatile esters, have pleasant, fruity smells and flavours. These three are components of the flavours of bananas, rum, and apples:



■
The terms 'saturated fats' and 'unsaturated fats' are familiar—they refer to whether the R groups are saturated (no C=C double bonds) or unsaturated (contains C=C double bonds)—see the box on p. 000. Fats containing R groups with several double bonds (for example, those that are esters formed from linoleic acid, which we met at the beginning of this chapter) are known as 'polyunsaturated'.

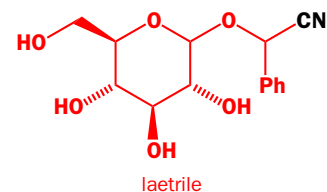
Amides (R-CONH₂, R¹-CONHR², or R¹CONR²R³)

Proteins are amides: they are formed when the carboxylic acid group of one amino acid condenses with the amino group of another to form an amide linkage (also known as a peptide bond). One protein molecule can contain hundreds of amide bonds. Aspartame, the artificial sweetener marketed as NutraSweet®, on the other hand contains just two amino acids, aspartic acid and phenylalanine, joined through one amide bond. Paracetamol is also an amide.



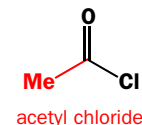
Nitriles or cyanides (R–CN) contain the cyano group –C≡N

Nitrile groups can be introduced into molecules by reacting potassium cyanide with alkyl halides. The organic nitrile group has quite different properties associated with lethal inorganic cyanide: Laetrile, for example, is extracted from apricot kernels, and was once developed as an anticancer drug. It was later proposed that the name be spelt ‘liar-trial’ since the results of the clinical trials on laetrile turned out to have been falsified!



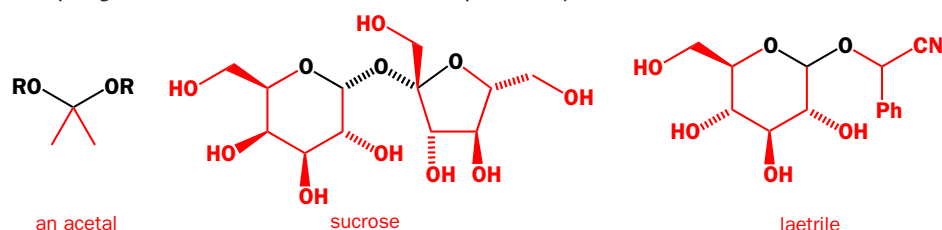
Acyl chlorides (acid chlorides)(R–COCl)

Acyl chlorides are reactive compounds used to make esters and amides. They are derivatives of carboxylic acids with the –OH replaced by –Cl, and are too reactive to be found in nature.



Acetals

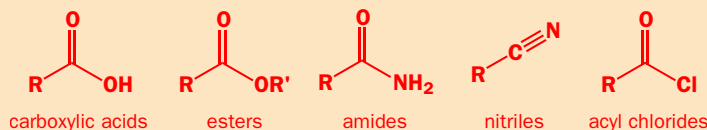
Acetals are compounds with two single bonded oxygen atoms attached to the same carbon atom. Many sugars are acetals, as is laetrile which you have just met.



Carbon atoms carrying functional groups can be classified by oxidation level

All functional groups are different, but some are more different than others. For example, the structures of a carboxylic acid, an ester, and an amide are all very similar: in each case the carbon atom carrying the functional group is bonded to two **heteroatoms**, one of the bonds being a double bond. You will see in Chapter 12 that this similarity in structure is mirrored in the reactions of these three types of compounds, and in the ways in which they can be interconverted. Carboxylic acids, esters, and amides can be changed one into another by reaction with simple reagents such as water, alcohols, or amines plus appropriate catalysts. To change them into aldehydes or alcohols requires a different type or reagent, a reducing agent (a reagent which adds hydrogen atoms). We say that the carbon atoms carrying functional groups that can be interconverted without the need for reducing agents (or oxidizing agents) have the same oxidation level—in this case, we call it the ‘carboxylic acid oxidation level’.

● The carboxylic acid oxidation level



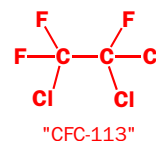
In fact, amides can quite easily be converted into nitriles just by dehydration (removal of water), so we must give nitrile carbon atoms the same oxidation level as carboxylic acids, esters, and amides. Maybe you’re beginning to see the structural similarity between these four functional groups that you could have used to assign their oxidation level? In all four cases, the carbon atom has *three* bonds to heteroatoms, and only one to C or H. It doesn’t matter how many heteroatoms there are, just how many bonds to them. Having noticed this, we can also assign both carbon atoms in ‘CFC-113’, one of the environmentally unfriendly aerosol propellants/refrigerants that have caused damage to the earth’s ozone layer, to the carboxylic acid oxidation level.

Aldehydes and ketones contain a carbon atom with *two* bonds to heteroatoms; they are at the ‘aldehyde oxidation level’. The common laboratory solvent dichloromethane also has two bonds to heteroatoms, so it too contains a carbon atom at the aldehyde oxidation level, as do acetals.

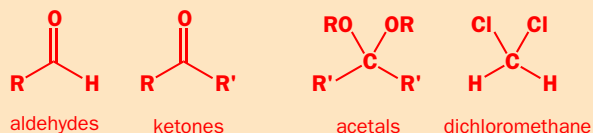
▶ A heteroatom is an atom that is not C or H

You’ve seen that a functional group is essentially any deviation from an alkane structure, either because the molecule has fewer hydrogen atoms than an alkane (alkenes, alkynes) or because it contains a collection of atoms that are not C and not H. There is a useful term for these ‘different’ atoms: heteroatoms. A **heteroatom** is any atom in an organic molecule other than C or H.

▶ Don’t confuse oxidation **level** with oxidation **state**. In all of these compounds, carbon is in oxidation state +4.

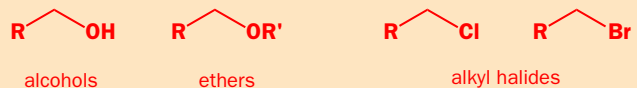


● The aldehyde oxidation level



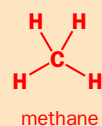
Alcohols, ethers, and alkyl halides have a carbon atom with only *one* single bond to a heteroatom. We assign these the 'alcohol oxidation level', and they are all easily made from alcohols without oxidation or reduction.

● The alcohol oxidation level



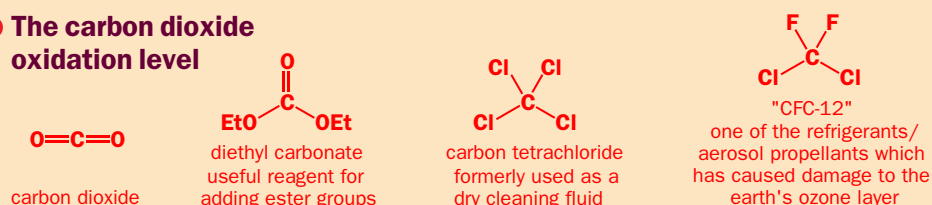
Lastly, we must include simple alkanes, which have no bonds to heteroatoms, as an 'alkane oxidation level'.

● The alkane oxidation level



The small class of compounds that have a carbon atom with four bonds to heteroatoms is related to CO₂ and best described as at the carbon dioxide oxidation level.

● The carbon dioxide oxidation level



● Summary: Important functional groups and oxidation levels

Zero bonds to heteroatoms Alkane oxidation level	One bond to heteroatom Alcohol oxidation level	Two bonds to heteroatoms Aldehyde oxidation level	Three bonds to heteroatoms Carboxylic acid oxidation level	Four bonds to heteroatoms Carbon dioxide oxidation level
 alkanes	alcohols ethers amines	aldehydes ketones acetals	carboxylic acids esters amides nitriles acyl chlorides	carbon dioxide diethyl carbonate CFC-12
	alkyl halides alkenes	alkynes		

Alkenes and alkynes obviously don't fit easily into these categories as they have no bonds to heteroatoms. Alkenes can be made from alcohols by dehydration without any oxidation or reduction so it seems sensible to put them in the alcohol column. Similarly, alkynes and aldehydes are related by hydration/dehydration without oxidation or reduction.

Naming compounds

So far, we have talked a lot about compounds by name. Many of the names we've used (palytoxin, muscone, brevetoxin...) are simple names given to complicated molecules without regard for the actual structure or function of the molecule—these three names, for example, are all derived from the name of the organism from which the compound was first extracted. They are known as **trivial names**, not because they are unimportant, but because they are used in everyday scientific conversation.

Names like this are fine for familiar compounds that are widely used and referred to by chemists, biologists, doctors, nurses, perfumers alike. But there are over 16 million known organic compounds. They can't all have simple names, and no one would remember them if they did. For this reason, the IUPAC (International Union of Pure and Applied Chemistry) have developed **systematic nomenclature**, a set of rules that allows any compound to be given a unique name that can be deduced directly from its chemical structure. Conversely, a chemical structure can be deduced from its systematic name.

The problem with systematic names is that they tend to be grotesquely unpronounceable for anything but the most simple molecules. In everyday speech and writing, chemists therefore do tend to disregard them, and use a mixture of systematic and trivial names. Nonetheless, it's important to know how the rules work. We shall look next at systematic nomenclature, before going on to look at the real language of chemistry.

Systematic nomenclature

There isn't space here to explain all the rules for giving systematic names for compounds—they fill several desperately dull volumes, and there's no point knowing them anyway since computers will do the naming for you. What we will do is to explain the principles underlying systematic nomenclature. You should understand these principles, because they provide the basis for the names used by chemists for the vast majority of compounds that do not have their own trivial names.

Systematic names can be divided into three parts: one describes the hydrocarbon framework; one describes the functional groups; and one indicates where the functional groups are attached to the skeleton.

You have already met the names for some simple fragments of hydrocarbon framework (methyl, ethyl, propyl...). Adding a hydrogen atom to these alkyl fragments and changing -yl to -ane makes the alkanes and their names. You should hardly need reminding of their structures:

Names for the hydrocarbon framework

one carbon methane **CH₄**



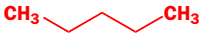








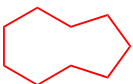

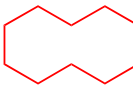
two carbons ethane **CH₃—CH₃**

three carbons propane **CH₃—CH₂—CH₃**

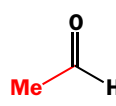
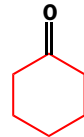
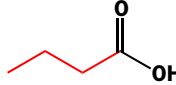
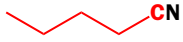
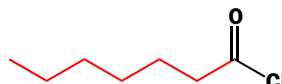
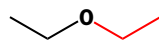

cyclopropane



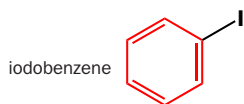
Names for the hydrocarbon framework (continued)

four carbons	butane		cyclobutane	
five carbons	pentane		cyclopentane	
six carbons	hexane		cyclohexane	
seven carbons	heptane		cycloheptane	
eight carbons	octane		cyclo-octane	
nine carbons	nonane		cyclononane	
ten carbons	decane		cyclodecane	

The name of a functional group can be added to the name of a hydrocarbon framework either as a suffix or as a prefix. Some examples follow. It is important to count all of the carbon atoms in the chain, even if one of them is part of a functional group: so pentanenitrile is actually BuCN.

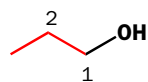
CH_3OH				
methanol	ethanal	cyclohexanone	butanoic acid	pentanenitrile
	$\text{HC}\equiv\text{CH}$		CH_3NO_2	
heptanoyl chloride	ethyne	ethoxyethane	nitromethane	propene

Compounds with functional groups attached to a benzene ring are named in a similar way.

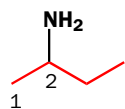


Numbers are used to locate functional groups

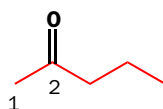
Sometimes a number can be included in the name to indicate which carbon atom the functional group is attached to. None of the above list needed a number—check that you can see why not for each one. When numbers are used, the carbon atoms are counted from one end. In most cases, either of two numbers could be used (depending on which end you count from); the one chosen is always the lower of the two. Again, some examples will illustrate this point. Notice again that some functional groups are named by prefixes, some by suffixes, and that the number always goes directly before the functional group name.



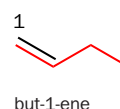
propan-1-ol



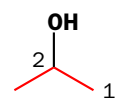
2-aminobutane



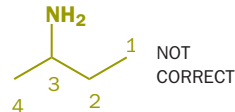
pentan-2-one



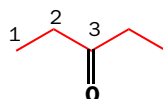
but-1-ene



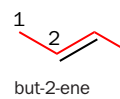
propan-2-ol



(not 3-aminobutane)

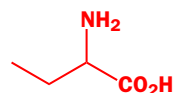


pentan-3-one



but-2-ene

Here are some examples of compounds with more than one functional group.



2-aminobutanoic acid



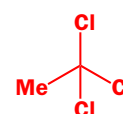
1,6-diaminohexane



hexanedioic acid



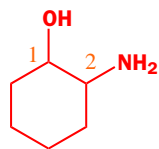
tetrabromomethane



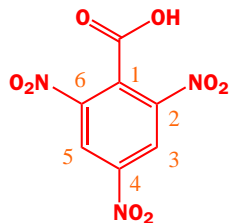
1,1,1-trichloroethane

Again, the numbers indicate how far the functional groups are from the end of the carbon chain. Counting must always be from the same end for each functional group. Notice how we use di-, tri-, tetra- if there are more than one of the same functional group.

With cyclic compounds, there isn't an end to the chain, but we can use numbers to show the distance between the two groups—start from the carbon atom carrying one of the functional groups, then count round.

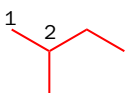


2-aminocyclohexanol

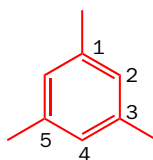


2,4,6-trinitrobenzoic acid

These rules work for hydrocarbon frameworks that are chains or rings, but many skeletons are branched. We can name these by treating the branch as though it were a functional group:



2-methylbutane



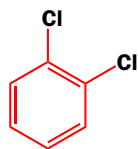
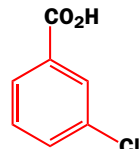
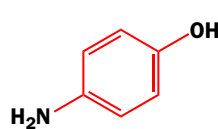
1,3,5-trimethyl benzene



1-butylcyclopropanol

Ortho, meta, and para

With substituted benzene rings, an alternative way of identifying the positions of the substituents is to use the terms *ortho*, *meta*, and *para*. *Ortho* compounds are 1,2-disubstituted, *meta* compounds are 1,3-disubstituted, and *para* compounds are 1,4-disubstituted. Some examples should make this clear.

1,2-dichlorobenzene
or *ortho*-dichlorobenzene
or *o*-dichlorobenzene3-chlorobenzoic acid
or *meta*-chlorobenzoic acid
or *m*-chlorobenzoic acid4-aminophenol
or *para*-aminophenol
or *p*-aminophenol

ortho, *meta*, and *para* are often abbreviated to *o*, *m*, and *p*.

▶ Beware! *Ortho*, *meta*, and *para* are used in chemistry to mean other things too: you may come across orthophosphoric acid, metastable states, and paraformaldehyde—these have nothing to do with the substitution patterns of benzene rings.

The terms *ortho*, *meta*, and *para* are used by chemists because they're easier to remember than numbers, and the words carry with them chemical meaning. '*Ortho*' shows that two groups are next to each other on the ring even though the atoms may not happen to be numbered 1 and 2. They are one example of the way in which chemists don't always use systematic nomenclature but revert to more convenient 'trivial' terms. We consider trivial names in the next section.

What do chemists really call compounds?

The point of naming a compound is to be able to communicate with other chemists. Most chemists are happiest communicating chemistry by means of structural diagrams, and structural drawings are far more important than any sort of chemical nomenclature. That's why we explained in detail how to draw structures, but only gave an outline of how to name compounds. Good diagrams are easy to understand, quick to draw, and difficult to misinterpret.

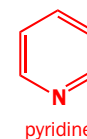
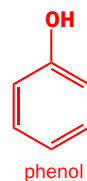
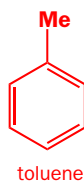
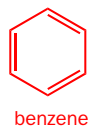
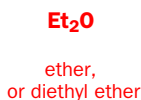
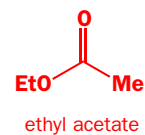
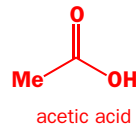
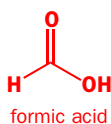
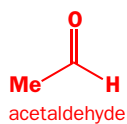
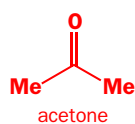
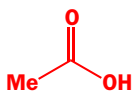
- Always give a diagram alongside a name unless it really is something very simple, such as ethanol.

But we do need to be able to communicate by speech and by writing as well. In principle we could do this by using systematic names. In practice, though, the full systematic names of anything but the simplest molecules are far too clumsy for use in everyday chemical speech. There are several alternatives, mostly based on a mixture of trivial and systematic names.

Names for well known and widely used simple compounds

A few simple compounds are called by trivial names not because the systematic names are complicated, but just out of habit. We know them so well that we use their familiar names.

You may have met this compound before (left), and perhaps called it ethanoic acid, its systematic name. But in a chemical laboratory, everyone would refer to this acid as acetic acid, its trivial name. The same is true for all these common substances.

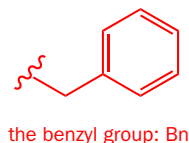
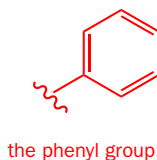
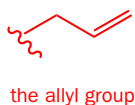
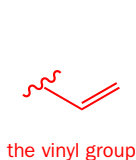


▶ We haven't asked you to remember any trivial names of molecules yet. But these 10 compounds are so important, you must be able to remember them. Learn them now.

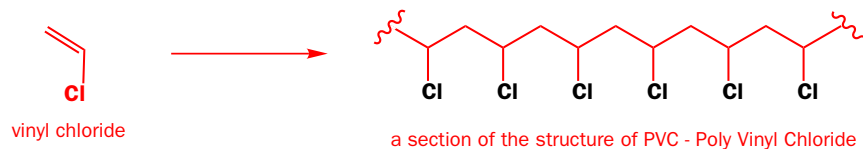
Trivial names like this are often long-lasting, well understood historical names that are less easy to confuse than their systematic counterparts. 'Acetaldehyde' is easier to distinguish from 'ethanol' than is 'ethanal'.

Trivial names also extend to fragments of structures containing functional groups. Acetone, acetaldehyde, and acetic acid all contain the acetyl group (MeCO-, ethanoyl) abbreviated Ac and chemists often use this 'organic element' in writing AcOH for acetic acid or EtOAc for ethyl acetate.

Chemists use special names for four fragments because they have mechanistic as well as structural significance. These are vinyl and allyl; phenyl and benzyl.



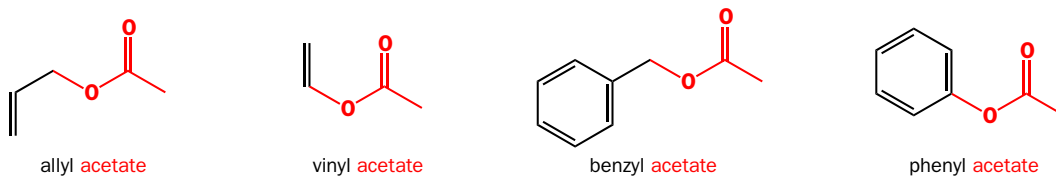
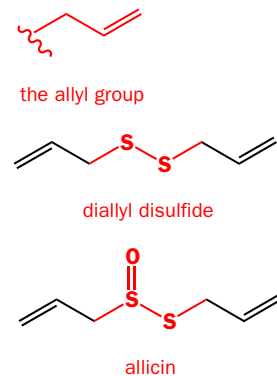
Giving the vinyl group a name allows chemists to use simple trivial names for compounds like vinyl chloride, the material that polymerizes to give PVC (poly vinyl chloride) but the importance of the name lies more in the difference in reactivity (Chapter 17) between vinyl and allyl groups.



The allyl group gets its name from garlic (*Allium* sp.), because it makes up part of the structure of the compounds responsible for the taste and smell of garlic.

Allyl and vinyl are different in that the vinyl group is attached directly to a double bonded C=C carbon atom, while the allyl group is attached to a carbon atom *adjacent* to the C=C double bond. The difference is extremely important chemically: allyl compounds are typically quite reactive, while vinyl compounds are fairly unreactive.

For some reason, the allyl and vinyl groups have never acquired organic element symbols, but the benzyl group has and is called Bn. It is again important not to confuse the benzyl group with the phenyl group: the phenyl group is joined through a carbon atom in the ring, while the benzyl group is joined through a carbon atom attached to the ring. Phenyl compounds are typically unreactive but benzyl compounds are often reactive. Phenyl is like vinyl and benzyl is like allyl.



We shall review all the organic element symbols you have met at the end of the chapter.

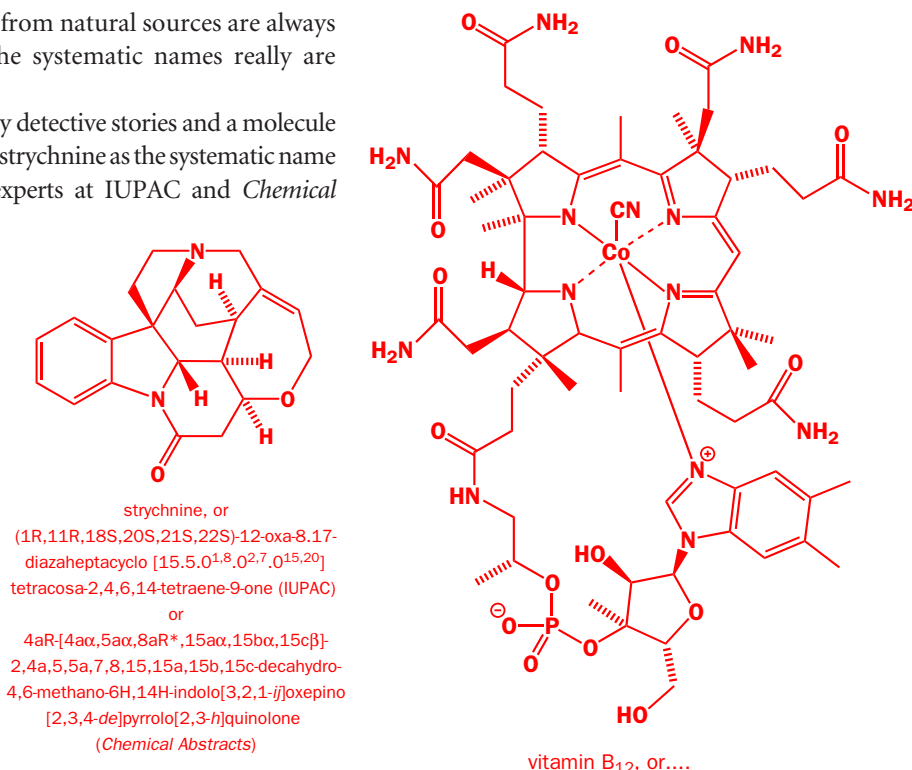
Names for more complicated but still well known molecules

Complicated molecules that have been isolated from natural sources are always given trivial names, because in these cases, the systematic names really are impossible!

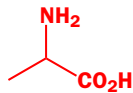
Strychnine is a famous poison featured in many detective stories and a molecule with a beautiful structure. All chemists refer to it as strychnine as the systematic name is virtually unpronounceable. Two groups of experts at IUPAC and *Chemical Abstracts* also have different ideas on the systematic name for strychnine. Others like this are penicillin, DNA, and folic acid.

But the champion is vitamin B₁₂, a complicated cobalt complex with a three-dimensional structure of great intricacy. No chemist would learn this structure but would look it up in an advanced textbook of organic chemistry. You will find it in such books in the index under vitamin B₁₂ and not under its systematic name. We do not even know what its systematic name might be and we are not very interested. This is vitamin B₁₂.

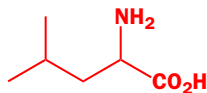
Even fairly simple but important molecules, the amino acids for example, that have systematic names that are relatively easy to understand are normally referred to by their



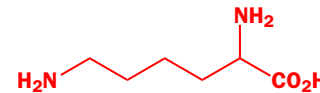
trivial names which are, with a bit of practice, easy to remember and hard to muddle up. They are given in full in Chapter 49.



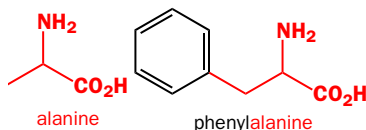
alanine, or
2-aminopropanoic acid



leucine, or
2-amino-4-methylpentanoic acid



lysine, or
2,6-diaminohexanoic acid

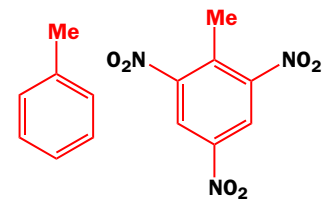


alanine

phenylalanine

A very flexible way of getting new, simple names for compounds can be to combine a bit of systematic nomenclature with trivial nomenclature.

Alanine is a simple amino acid that occurs in proteins. Add a phenyl group and you have phenylalanine a more complex amino acid also in proteins.



toluene

2,4,6-trinitrotoluene

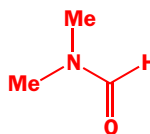
Toluene, the common name for methylbenzene, can be combined (both chemically and in making names for compounds!) with three nitro groups to give the famous explosive trinitrotoluene or TNT.

Compounds named as acronyms

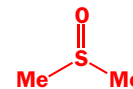
Some compounds are referred to by acronyms, shortened versions of either their systematic or their trivial name. We just saw TNT as an abbreviation for TriNitroToluene but the commoner use for acronyms is to define solvents and reagents in use all the time. Later in the book you will meet these solvents.



THF
(TetraHydroFuran)



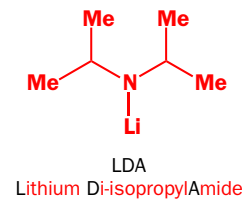
DMF
(DiMethylFormamide)



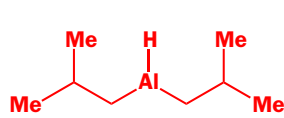
DMSO
(DiMethylSulfoxide)

▶ The names and structures of these common solvents need learning too.

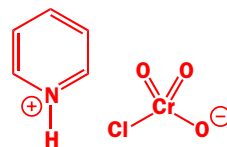
The following reagents are usually referred to by acronym and their functions will be introduced in other chapters so you do not need to learn them now. You may notice that some acronyms refer to trivial and some to systematic names. There is a glossary of acronyms for solvents, reagents, and other compounds on p. 000.



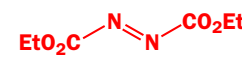
LDA
Lithium Di-isopropylAmide



DIBAL
Di-IsoButylAluminiumhydride



PCC
Pyridinium ChloroChromate



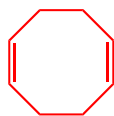
DEAD
DiEthyl Azo-Dicarboxylate

Compounds for which chemists use systematic names

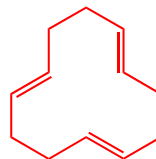
You may be surprised to hear that practising organic chemists use systematic names at all in view of what we have just described, but they do! Systematic names really begin with derivatives of pentane (C_5H_{12}) since the prefix pent- means five, whereas but- does not mean four. Chemists refer to simple derivatives of open chain and cyclic compounds with 5 to about 20 carbon atoms by their systematic names, providing that there is no common name in use. Here are some examples.



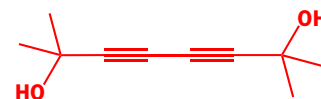
cyclopentadiene



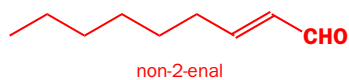
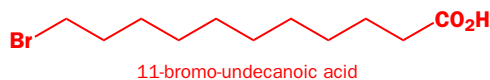
cyclo-octa-1,5-diene



cyclododeca-1,5,9-triene



2,7-dimethyl-3,5-octadiyne-2,7-diol

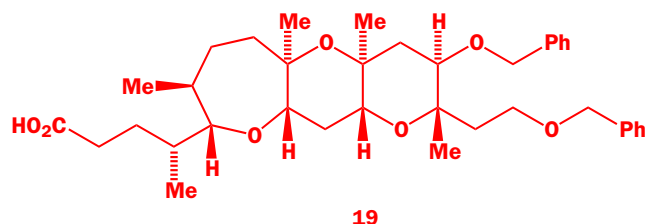


These names contain a syllable that tells you the framework size: penta- for C₅, octa- for C₈, nona- for C₉, undeca- for C₁₁, and dodeca- for C₁₂. These names are easily worked out from the structures and, what is more important, you get a clear idea of the structure from the name. One of them might make you stop and think a bit (which one?), but the others are clear even when heard without a diagram to look at.

Complicated molecules with no trivial names

When chemists make complex new compounds in the laboratory, they publish them in a chemical journal giving their full systematic names in the experimental account, however long and clumsy those names may be. But in the text of the paper, and while talking in the lab about the compounds they have made, they will just call them ‘the amine’ or ‘the alkene’. Everyone knows which amine or alkene is meant because at some point they remember seeing a chemical structure of the compound. This is the best strategy for talking about almost any molecule: draw a structure, then give the compound a ‘tag’ name like ‘the amine’ or ‘the acid’. In written chemistry it’s often easiest to give every chemical structure a ‘tag’ number as well.

To illustrate what we mean, let’s talk about this compound.



This carboxylic acid was made and used as an intermediate when chemists in California made brevetoxin (see p. 000) in 1995. Notice how we can call a complicated molecule ‘this acid’—a ‘tag’ name—because you’ve seen the structure. It also has a tag number (19), so we can also call it ‘compound 19’, or ‘acid 19’, or ‘brevetoxin fragment 19’. How much more sensible than trying to work out its systematic name.

How should you name compounds?

So what should you call a compound? It really depends on circumstances, but you won’t go far wrong if you follow the example of this book. We shall use the names for compounds that real

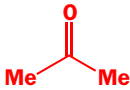
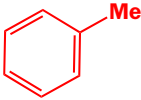
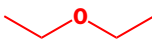

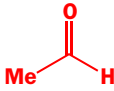
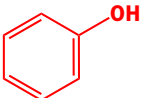
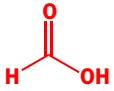

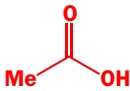
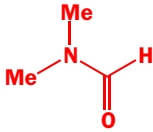

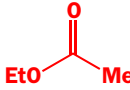
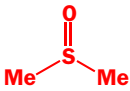
● Our advice on chemical names—six points in order of importance

- Draw a structure first and worry about the name afterwards
- Learn the names of the *functional groups* (ester, nitrile, etc.)
- Learn and use the names of a few simple compounds used by all chemists
- In speech, refer to compounds as ‘that acid’ (or whatever) while pointing to a diagram
- Grasp the principles of systematic (IUPAC) nomenclature and use it for compounds of medium size
- Keep a notebook to record acronyms, trivial names, structures, etc. that you might need later

chemists use. There's no need to learn all the commonly used names for compounds now, but you should log them in your memory as you come across them. Never allow yourself to pass a compound name by unless you are sure you know what chemical structure it refers to. You will find many of the commonly used names for compounds on the endpapers of this book. Refer to these, or to the short-term glossary on p. 000 to refresh your memory should you ever need to.




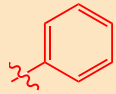

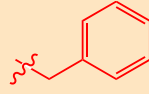




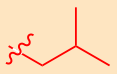
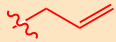
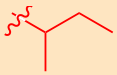
We've met a great many molecules in this chapter. Most of them were just there to illustrate points so don't learn their structures! Instead, learn to recognize the names of the functional groups they contain. However, there were 10 names for simple compounds and three for common solvents that we advised you to learn. Cover up the structures on the rest of this page and draw the structures for these 13 compounds.

Important structures to learn

acetone		toluene	
ether or diethyl ether		pyridine	
acetaldehyde		phenol	
formic acid		THF (tetrahydrofuran)	
acetic acid or AcOH		DMF or (dimethylformamide) Me ₂ NCHO	
benzene			
ethyl acetate or EtOAc		DMSO (dimethylsulfoxide)	

That's all we'll say on the subject of nomenclature—you'll find that as you practise using these names and start hearing other people referring to compounds by name you'll soon pick up the most important ones. But, to reiterate, make sure you never pass a compound name by without being absolutely sure what it refers to—draw a structure to check.

● Review box: Table of fragment names and organic elements

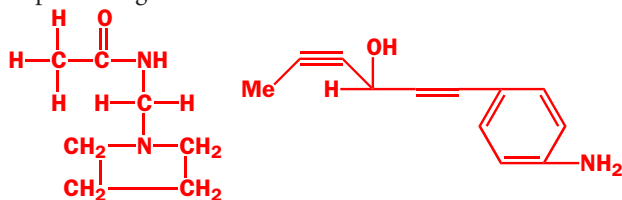
R	alkyl		t-Bu	<i>tert</i> -butyl	
Me	methyl		Ar	aryl	any aromatic ring
Et	ethyl		Ph	phenyl	
Pr (or n-Pr)	propyl		Bn	benzyl	
Bu (or n-Bu)	butyl		Ac	acetyl	
i-Pr	isopropyl			vinyl	
i-Bu	isobutyl			allyl	
s-Bu	sec-butyl		X	halide	F, Cl, Br, or I

Problems

1. Draw good diagrams of saturated hydrocarbons with seven carbon atoms having (a) linear, (b) branched, and (c) cyclic frameworks. Draw molecules based on each framework having both ketone and carboxylic acid functional groups.

2. Study the structure of brevetoxin on p. 000. Make a list of the different types of functional group (you already know that there are many ethers) and of the numbers of rings of different sizes. Finally study the carbon framework—is it linear, cyclic, or branched?

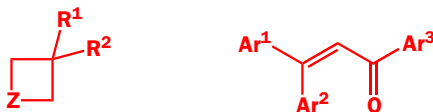
3. What is wrong with these structures? Suggest better ways of representing these molecules.



4. Draw structures corresponding to these names. In each case suggest alternative names that might convey the structure more clearly to someone who is listening to you speak.

- 1,4-di-1(1-dimethylethyl)benzene
- 2-(prop-2-enyloxy)prop-1-ene
- cyclohexa-1,3,5-triene

5. Draw one possible structure for each of these molecules, selecting any group of your choice for the 'wild card' substituents.



6. Translate these very poor 'diagrams' of molecules into more realistic structures. Try to get the angles about right and, whatever you do, don't include any square coplanar carbon atoms or other bond angles of 90°!



7. Suggest at least six different structures that would fit the formula $\text{C}_4\text{H}_7\text{NO}$. Make good realistic diagrams of each one and say which functional group(s) are present.

8. Draw and name a structure corresponding to each of these descriptions.

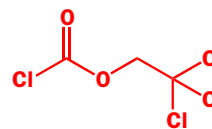
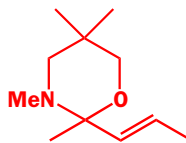
- An aromatic compound containing one benzene ring with the following substituents: two chlorine atoms having a *para* relationship, a nitro group having an *ortho* relationship to one of the chlorine atoms, and an acetyl group having a *meta* relationship to the nitro group.

(b) An alkyne having a trifluoromethyl substituent at one end and a chain of three carbon atoms at the other with a hydroxyl group on the first atom, an amino group on the second, and the third being a carboxyl group.

9. Draw full structures for these compounds, displaying the hydrocarbon framework clearly and showing all the bonds present in the functional groups. Name the functional groups.



10. Identify the oxidation level of each of the carbon atoms in these structures with some sort of justification.



11. If you have not already done so, complete the exercises on pp. 000 (drawing amino acids) and 000 (giving structures for the 10 common compounds and three common solvents).