Learning Objectives

- To understand how catalysts increase the reaction rate and the selectivity of chemical reactions.

Catalysts are substances that increase the reaction rate of a chemical reaction without being consumed in the process. A catalyst, therefore, does not appear in the overall stoichiometry of the reaction it catalyzes, but it must appear in at least one of the elementary reactions in the mechanism for the catalyzed reaction. The catalyzed pathway has a lower $E_a$, but the net change in energy that results from the reaction (the difference between the energy of the reactants and the energy of the products) is not affected by the presence of a catalyst (Figure \(\PageIndex{1}\)). Nevertheless, because of its lower $E_a$, the reaction rate of a catalyzed reaction is faster than the reaction rate of the uncatalyzed reaction at the same temperature. Because a catalyst decreases the height of the energy barrier, its presence increases the reaction rates of both the forward and the reverse reactions by the same amount. In this section, we will examine the three major classes of catalysts: heterogeneous catalysts, homogeneous catalysts, and enzymes.

![Potential energy diagrams for a single-step reaction in the presence and absence of a catalyst](image1)

**Figure \(\PageIndex{1}\): Lowering the Activation Energy of a Reaction by a Catalyst.** This graph compares potential energy diagrams for a single-step reaction in the presence and absence of a catalyst. The only effect of the catalyst is to lower the activation energy of the reaction. The catalyst does not affect the energy of the reactants or products (and thus does not affect $\Delta E$).

A catalyst affects $E_a$, not $\Delta E$.

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**Heterogeneous Catalysis**

In **heterogeneous catalysis**, the catalyst is in a different phase from the reactants. At least one of the reactants interacts with the solid surface in a physical process called adsorption in such a way that a chemical bond in the reactant becomes weak and then breaks. Poisons are substances that bind irreversibly to catalysts, preventing reactants from adsorbing and thus reducing or destroying the catalyst’s efficiency.

An example of heterogenous catalysis is the interaction of hydrogen gas with the surface of a metal, such as Ni, Pd, or Pt. As shown in part (a) in Figure \(\PageIndex{2}\), the hydrogen–hydrogen bonds break and produce individual adsorbed hydrogen atoms on the surface of the metal. Because the adsorbed atoms can move around on the surface, two hydrogen atoms can collide and form a molecule of hydrogen gas that can then leave the surface in the reverse process, called desorption. Adsorbed H atoms on a metal surface are substantially more reactive than a hydrogen...
molecule. Because the relatively strong H–H bond (dissociation energy = 432 kJ/mol) has already been broken, the
energy barrier for most reactions of H₂ is substantially lower on the catalyst surface.

Figure \(\PageIndex{2}\): Hydrogenation of Ethylene on a Heterogeneous Catalyst. When a molecule of hydrogen
adsorbs to the catalyst surface, the H–H bond breaks, and new M–H bonds are formed. The individual H atoms are
more reactive than gaseous H₂. When a molecule of ethylene interacts with the catalyst surface, it reacts with the H
atoms in a stepwise process to eventually produce ethane, which is released.

Figure \(\PageIndex{2}\) shows a process called hydrogenation, in which hydrogen atoms are added to the double bond
of an alkene, such as ethylene, to give a product that contains C–C single bonds, in this case ethane. Hydrogenation is
used in the food industry to convert vegetable oils, which consist of long chains of alkenes, to more commercially
valuable solid derivatives that contain alkyl chains. Hydrogenation of some of the double bonds in polyunsaturated
vegetable oils, for example, produces margarine, a product with a melting point, texture, and other physical properties
similar to those of butter.

Several important examples of industrial heterogeneous catalytic reactions are in Table \(\PageIndex{1}\). Although the
mechanisms of these reactions are considerably more complex than the simple hydrogenation reaction described here,
they all involve adsorption of the reactants onto a solid catalytic surface, chemical reaction of the adsorbed species
(sometimes via a number of intermediate species), and finally desorption of the products from the surface.

<table>
<thead>
<tr>
<th>Commercial Process</th>
<th>Catalyst</th>
<th>Initial Reaction</th>
</tr>
</thead>
<tbody>
<tr>
<td>contact process</td>
<td>V₂O₅ or Pt</td>
<td>2SO₂ + O₂ → 2SO₃</td>
</tr>
<tr>
<td>Haber process</td>
<td>Fe, K₂O, Al₂O₃</td>
<td>N₂ + 3H₂ → 2NH₃</td>
</tr>
<tr>
<td>Ostwald process</td>
<td>Pt and Rh</td>
<td>4NH₃ + 5O₂ → 4NO + 6H₂O</td>
</tr>
<tr>
<td>water–gas shift reaction</td>
<td>Fe, Cr₂O₃, or Cu</td>
<td>CO + H₂O → CO₂ + H₂</td>
</tr>
</tbody>
</table>
Commercial Process | Catalyst | Initial Reaction
--- | --- | ---
steam reforming | Ni | CH₄ + H₂O → CO + 3H₂
methanol synthesis | ZnO and Cr₂O₃ | CO + 2H₂ → CH₃OH
Sohio process | bismuth phosphomolybdate | CO + 2H₂ → CH₃OH
Catalytic hydrogenation | Ni, Pd, or Pt | RCH=CHR' + H₂ → RCH₂—CH₂R'

Homogeneous Catalysis

In **homogeneous catalysis**, the catalyst is in the same phase as the reactant(s). The number of collisions between reactants and catalyst is at a maximum because the catalyst is uniformly dispersed throughout the reaction mixture. Many homogeneous catalysts in industry are transition metal compounds (Table \(\PageIndex{2}\)), but recovering these expensive catalysts from solution has been a major challenge. As an added barrier to their widespread commercial use, many homogeneous catalysts can be used only at relatively low temperatures, and even then they tend to decompose slowly in solution. Despite these problems, a number of commercially viable processes have been developed in recent years. High-density polyethylene and polypropylene are produced by homogeneous catalysis.

Table \(\PageIndex{2}\): Some Commercially Important Reactions that Employ Homogeneous Catalysts

<table>
<thead>
<tr>
<th>Commercial Process</th>
<th>Catalyst</th>
<th>Reactants</th>
<th>Final Product</th>
</tr>
</thead>
<tbody>
<tr>
<td>Union Carbide</td>
<td>[Rh(CO)₂I₂]⁻</td>
<td>CO + CH₃OH</td>
<td>CH₃CO₂H</td>
</tr>
<tr>
<td>Hydroperoxide process</td>
<td>Mo(VI) complexes</td>
<td>CH₃CH=CH₂ + R–O–O–H</td>
<td>Propylene oxide</td>
</tr>
<tr>
<td>Hydroformylation</td>
<td>Rh/PR₃ complexes</td>
<td>RCH=CH₂ + CO + H₂</td>
<td>RCH₂CH₂CHO</td>
</tr>
<tr>
<td>Adiponitrile process</td>
<td>Ni/PR₃ complexes</td>
<td>2HCN + CH₂=CHCH=CH₂</td>
<td>NCCH₂CH₂CH₂CH₂CN used to synthesize nylon</td>
</tr>
</tbody>
</table>
### Enzymes

Enzymes, catalysts that occur naturally in living organisms, are almost all protein molecules with typical molecular masses of 20,000–100,000 amu. Some are homogeneous catalysts that react in aqueous solution within a cellular compartment of an organism. Others are heterogeneous catalysts embedded within the membranes that separate cells and cellular compartments from their surroundings. The reactant in an enzyme-catalyzed reaction is called a substrate.

Because enzymes can increase reaction rates by enormous factors (up to $10^{17}$ times the uncatalyzed rate) and tend to be very specific, typically producing only a single product in quantitative yield, they are the focus of active research. At the same time, enzymes are usually expensive to obtain, they often cease functioning at temperatures greater than 37 °C, have limited stability in solution, and have such high specificity that they are confined to turning one particular set of reactants into one particular product. This means that separate processes using different enzymes must be developed for chemically similar reactions, which is time-consuming and expensive. Thus far, enzymes have found only limited industrial applications, although they are used as ingredients in laundry detergents, contact lens cleaners, and meat tenderizers. The enzymes in these applications tend to be proteases, which are able to cleave the amide bonds that hold amino acids together in proteins. Meat tenderizers, for example, contain a protease called papain, which is isolated from papaya juice. It cleaves some of the long, fibrous protein molecules that make inexpensive cuts of beef tough, producing a piece of meat that is more tender. Some insects, like the bombadier beetle, carry an enzyme capable of catalyzing the decomposition of hydrogen peroxide to water (Figure \(\PageIndex{3}\)).

![A Catalytic Defense Mechanism](image)

**Figure \(\PageIndex{3}\):** A Catalytic Defense Mechanism. The scalding, foul-smelling spray emitted by this bombardier beetle is produced by the catalytic decomposition of \(\ce{H2O2}\).

Enzyme inhibitors cause a decrease in the reaction rate of an enzyme-catalyzed reaction by binding to a specific portion of an enzyme and thus slowing or preventing a reaction from occurring. Irreversible inhibitors are therefore the equivalent of poisons in heterogeneous catalysis. One of the oldest and most widely used commercial enzyme inhibitors is aspirin, which selectively inhibits one of the enzymes involved in the synthesis of molecules that trigger inflammation.
The design and synthesis of related molecules that are more effective, more selective, and less toxic than aspirin are important objectives of biomedical research.

**Summary**

Catalysts participate in a chemical reaction and increase its rate. They do not appear in the reaction’s net equation and are not consumed during the reaction. Catalysts allow a reaction to proceed via a pathway that has a lower activation energy than the uncatalyzed reaction. In heterogeneous catalysis, catalysts provide a surface to which reactants bind in a process of adsorption. In homogeneous catalysis, catalysts are in the same phase as the reactants. Enzymes are biological catalysts that produce large increases in reaction rates and tend to be specific for certain reactants and products. The reactant in an enzyme-catalyzed reaction is called a substrate. Enzyme inhibitors cause a decrease in the reaction rate of an enzyme-catalyzed reaction.