

The previous equation assumes slow binding of enzyme (i.e.,  $[E] \approx [E_0]$ ),  $S_0$  is the number of substrate bonds available initially for breakage, and  $k_{\text{des}}$  and  $k_{\text{ads}}$  refer to rates of enzyme desorption and adsorption onto the insoluble matrix, respectively.

### 3.4 IMMOBILIZED ENZYME SYSTEMS

The restriction of enzyme mobility in a fixed space is known as *enzyme immobilization*. Immobilization of enzymes provides important advantages, such as enzyme reutilization and elimination of enzyme recovery and purification processes, and may provide a better environment for enzyme activity. Since enzymes are expensive, catalyst reuse is critical for many processes. Since some of the intracellular enzymes are membrane bound, immobilized enzymes provide a model system to mimic and understand the action of some membrane-bound intracellular enzymes. Product purity is usually improved, and effluent handling problems are minimized by immobilization.

#### 3.4.1. Methods of Immobilization

Major methods of immobilization are summarized in Fig. 3.16. The two major categories are entrapment and surface immobilization.

**3.4.1.1. Entrapment.** Entrapment is the physical enclosure of enzymes in a small space. Matrix entrapment and membrane entrapment, including microencapsulation, are the two major methods of entrapment.

Matrices used for enzyme immobilization are usually polymeric materials such as Ca-alginate, agar,  $\kappa$ -carrageenin, polyacrylamide, and collagen. However, some solid matrices such as activated carbon, porous ceramic, and diatomaceous earth can also be used for this purpose. The matrix can be a particle, a membrane, or a fiber. When immobilizing

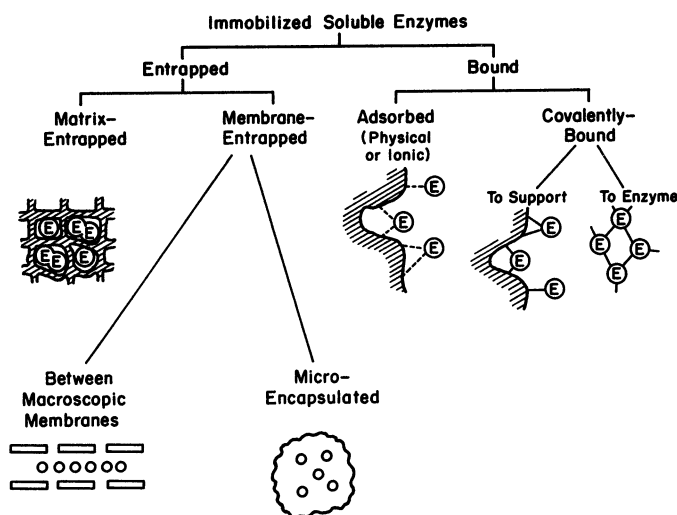


Figure 3.16. Major immobilization methods.