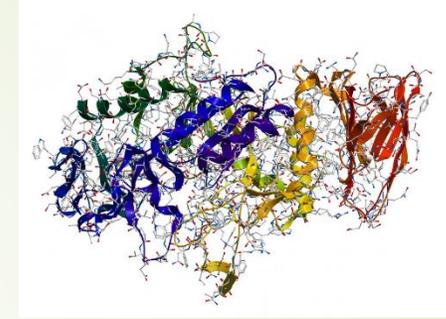


# CEN-CHE 422

## ENZYME ENGINEERING



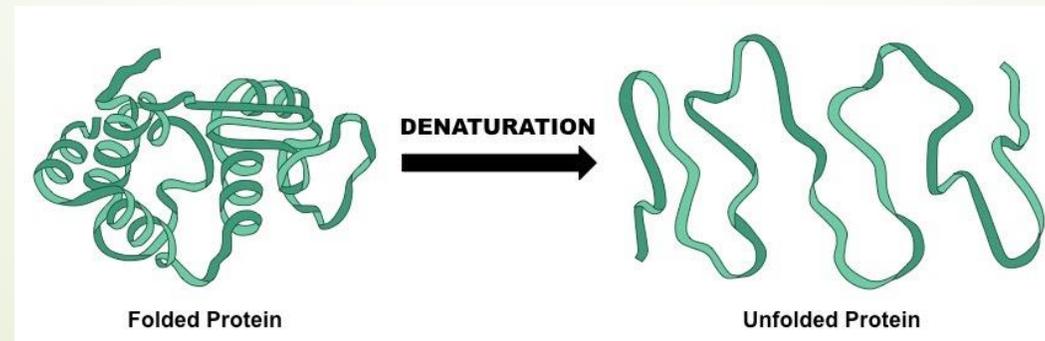
# ENZYME DEACTIVATION AND STABILITY

2

**Stability:** Capacity of the enzyme to retain its catalytic activity

**Physical, chemical and biochemical factors** (pH, temperature, additives)

**Deactivation:** Structural and catalytic



## Physical Deactivation:

High temperature

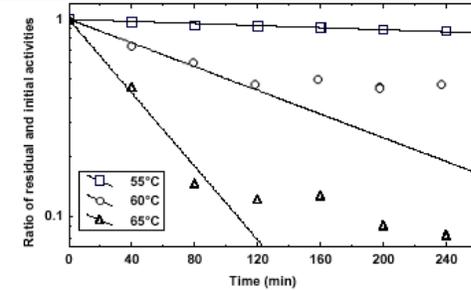
Low temperature

pH

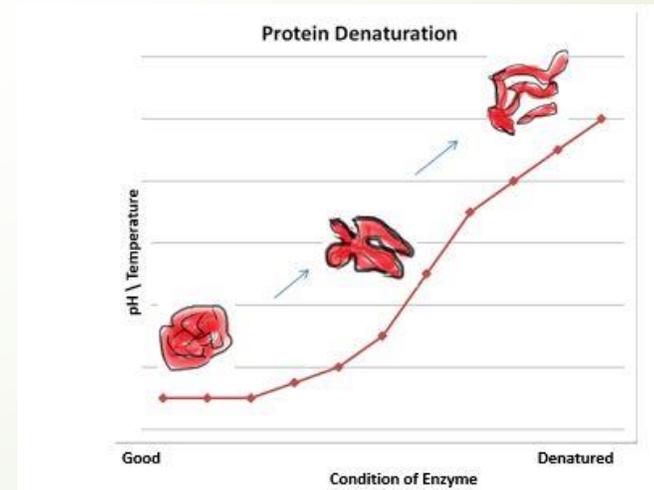
Mechanical forces

## Biological inactivation :

Protease



**Figure 2:** Immobilized invertase thermal deactivation data compared with the exponential decay model. One gram of the immobilized enzyme, containing 15.75 mg protein/g of support was incubated in 200 mL of substrate solution containing 5% (w/v) sucrose buffered with disodium phosphate-citric acid buffer 50 mM, at pH 4.5.



## Chemical deactivation:

Organic H-bond forming

Oxidants

Neutral salts :

Solvents

## Stability

1. Genetic and protein engineering
2. Medium Engineering
3. Immobilization
4. CLEC
5. Additives
6. Chemical modification