

Name \_\_\_\_\_ ID # \_\_\_\_\_

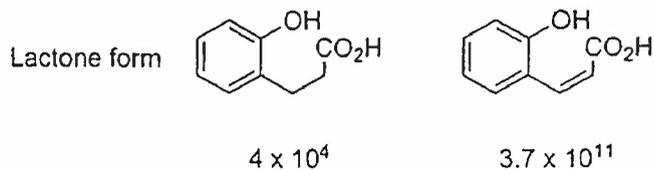
SB  
2019/2018

1. What is the primary drawback of the lock-and-key model in explaining catalysis? Use an appropriate diagram to answer. Lengthy discussion is not encouraged. [4]

*Diagram :**Explanation :*

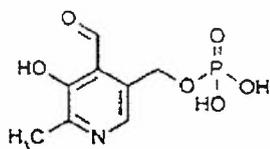
2. In the report of an enzyme kinetics experiment, a student reported an experimentally obtained second-order rate constant of  $1.5 \times 10^{17} \text{ M}^{-1} \text{ s}^{-1}$  with an enzyme (5  $\mu\text{M}$ ) and its natural substrate (1, 2, 5, 10, 25, 50 mM). Does the result look credible to you? Explain. [4]

3. (a) The rate constants (in  $\text{h}^{-1}$ ) of lactonization (lactone = cyclic ester) for the following substrates are provided with their structures. Why does the second substrate react faster? [2]



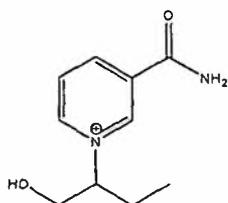
- (b) To calculate the effective molarity from these rate constants provided above, what other substrate(s) would be required for comparison? Write the reaction. [2]

4. Alanine does not racemize in  $\text{H}_2\text{O}$  under physiological pH, however in the presence of PLP (structure given below), it can. Why? Write a relevant structure to explain your point. [3]



5. Normally, enzyme catalysis is achieved by *stabilization* of the transition state of a reaction. Can a *destabilizing factor* be employed to enhance the reaction rate in a catalysis? Draw it in an energy profile diagram to demonstrate. You do not need to write too much. A structure or figure says a thousand words. [3]

6. If the following molecule binds to an antibody effectively, show **three** different types of interactions between the molecule and the amino acid side chains (structure required) of the antibody. [2]



**2018 Midsem Exam ID 4107 (ANSWER WITHIN THE GIVEN SPACE)**

**Space for rough work**