## Chemistry 112B: Midterm 1, Tuesday February 21, 2017

Name:	

UCSID:	<i>GSI</i> :
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There are a total of <u>11 pages</u> on this exam including this one. Time for the exam **8:10AM –9:30 AM**. By writing your name on this exam, you have acknowledged that you have all 11 pages and written your answers only on the designated pages with page numbers (*answers written on the back pages will NOT be graded*).

Question 1 \_\_\_\_\_(17 pts)

Question 2 \_\_\_\_\_(10 pts)

Question 3 \_\_\_\_\_(10 pts)

Question 4 \_\_\_\_\_(12 pts)

Question 5 \_\_\_\_\_(14 pts)

Question 6 \_\_\_\_\_(12 pts)

Question 7 \_\_\_\_\_(10 pts)

Question 8 \_\_\_\_\_(15 pts)

Total \_\_\_\_/100 points

(a) List the names of the following functional groups on the lines (1 pt each)



(b) Rank the following functional groups from 1-5 with the most electrophilic as 1 (1pt each)



A) Circle the diagram  $(\mathbf{A}, \mathbf{B}, \text{ or } \mathbf{C})$  that best represents the approach of a nucleophile to form a tetrahedral intermediate (2 pts)



B) Provide a rationalization for your answer in Part A in no more than 2 sentences and 2 figures. (8 pts)

Question 3





Fill in the reagents or products in the boxes that are provided. (3 points each)

(a)









(a) Provide a base (**X**) that will lead to the *reversible* deprotonation of **D** (indicated with equilibrium arrows) or a base (**Y**) that would lead to *irreversible* deprotonation of **D** (1 pt each).



(b) Explain with one figure and in three sentences or less why deprotonation of **D** with **Y** in Part(a) leads to irreversible deprotonation (4 pts) (**Hint:** Consider the conjugate acid that is formed).

(c) Provide a rationalization for the following observations using base **Y** from Part (a) (or any base that gives *kinetic*, *irreversible* deprotonation). You may use up to four figures and five sentences. (8 pts)



Fill in the blank boxes below with the required compounds, reagents, or names.

### A) 2 pts each



B) 2 pts each



C) 2 pts each



(a) Provide conditions and the name associated with the conversion of  $\mathbf{Q}$  to  $\mathbf{R}$  shown below. (4 pts + 2 pts)



(b) Why isn't **X** (below) the final product in the reaction of **Q** under the conditions you listed above? Explain in three (or less) sentences and three (or less) structures (4 pts)



Amide bond forming reactions are used regularly in the pharmaceutical industry to make many medicines including the pain medication fentanyl. Fentanyl may be prepared from amine **XX** and acid **YY** through a DCC coupling.



A) Provide the structures of two <u>different</u> "active ester" functional groups that can be prepared from **YY** separate from the one that is generated from DCC. (3 pts each)





B) <u>One the following page</u>, provide a detailed mechanism for the formation of fentanyl from **XX** and **YY** using DCC. (9 points)

# The End