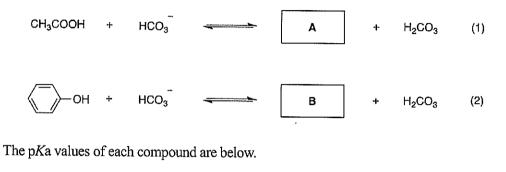
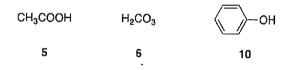
## Organic Chemistry I

I - a

Answer the following questions in the reaction equilibrium.





- (1) Write the structure of the A and B.
- (2) Regarding acetic acid and carbon acid in equation (1), which one is a stronger acid? Show how (what times) strong it is based on the pKa value.
- (3) In equation (2), does the reaction equilibrium lie to the right or the left? Explain the reason based on the pKa value?
- I b

Predict the major product and discuss the basis for the stereochemistry resulting from reaction of 1-methylcyclohexene with Br<sub>2</sub>.

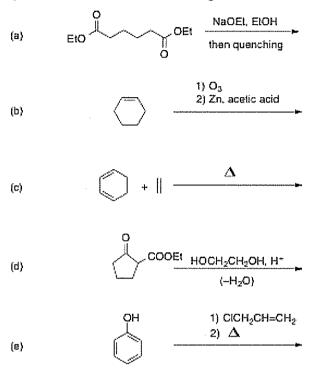
I — c

Predict the product and discuss the basis for the stereochemistry resulting from reaction of each of the following reagents with (S)-2-bromohexane.

(1) NaI

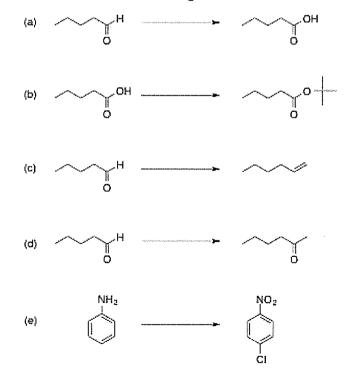
(2) KCN

Give the structure of the product obtained from the following reactions.



I - e

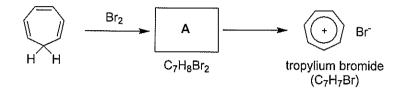
Give the reagents and/or conditions for the following transformations.



## Organic Chemistry II

II — a

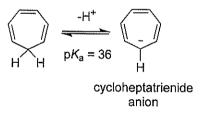
(1) Cycloheptatriene reacts with bromine to give compound A. Draw the structure of A.



(2) The distillation of compound A gave tropylium bromide as the yellow solid. Draw the reaction scheme.

(3) How many  $\pi$  electrons does tropylium ion have? Why is the ionic form more stable than the neutral form?

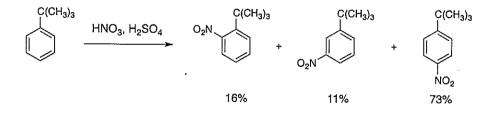
(4) The following show the p $K_a$  of cycloheptatriene. How many  $\pi$  electrons does its anion have?



(5) Why is cycloheptatrienide anion far less stable than tropylium ion?

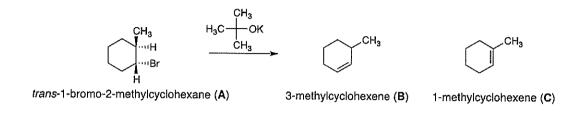
 $\rm I\!I-b$ 

Nitration of *t*-butylbenzene affords *ortho-*, *meta-*, and *para-*substituted products with the indicated selectivity. Explain the reason for this selectivity with the reaction mechanism.



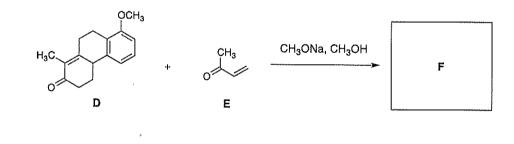
 $\rm I\!I-c$ 

Reaction between *trans*-1-bromo-2-methylcyclohexane (A) and potassium *t*-butoxide affords methylcyclohexene through E2 *anti*-elimination mechanism. Answer which is the product, 3-methylcyclohexene (B) or 1-methylcyclohexene (C). And explain the reason with the reaction mechanism.



## $\Pi - d$

Compounds **D** and **E** undergo Michael addition and aldol condensation reactions to afford product **F**. Answer the structure of compound **F** and the reaction mechanism.



 $\Pi - e$ 

This scheme shows the synthesis of (S)-Actinidiolide. Answer the structure of product G.

