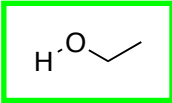
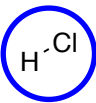
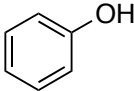
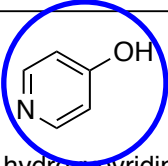
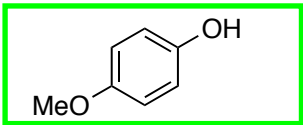
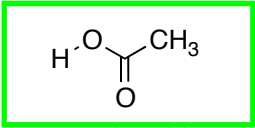
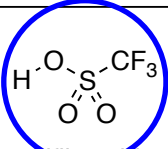
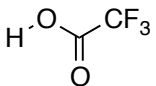
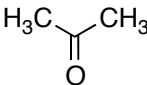
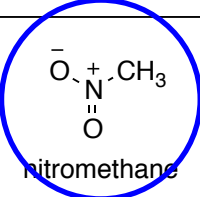
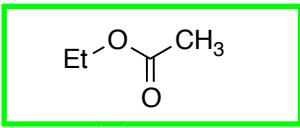
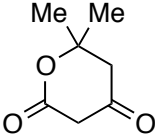
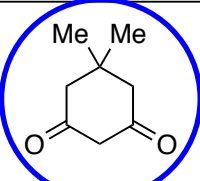
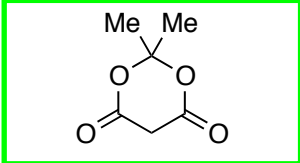


Name _____ **Answer Key** _____Part I 10 / 10Part II 18 / 18Part III 24 / 24Part IV 20 / 20Part V 8 / 08Part VI 20 / 20Total 100 / 100

There are 100 points on the exam.

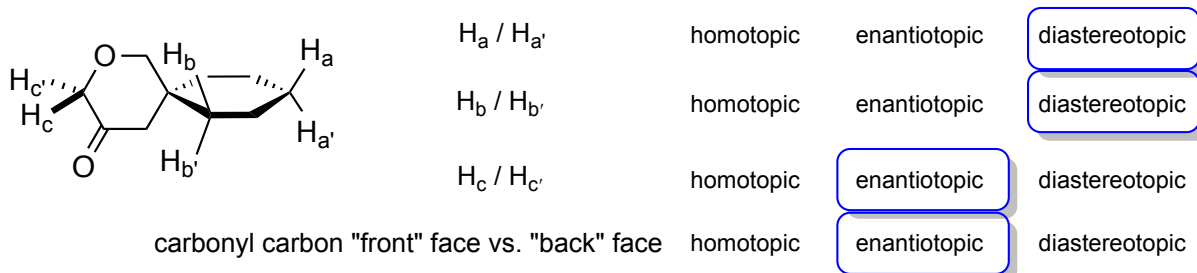
Answer all six of questions I-VI directly in the space provided on the six exam pages. You may not use books, notes, models, etc.

I. (10 points) Within each set, **circle the strongest acid** and place a rectangle around the weakest acid.

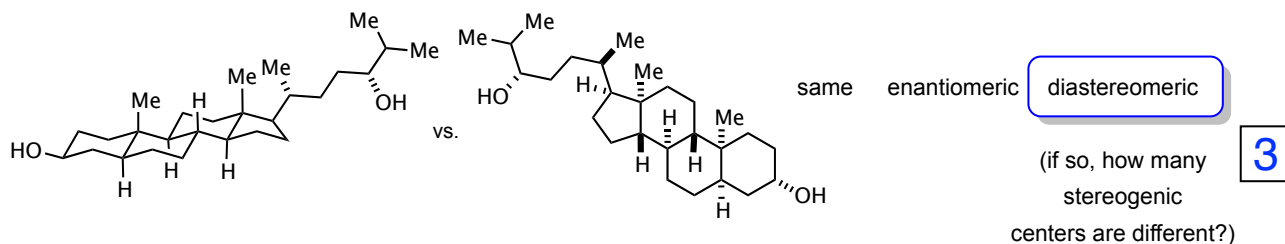
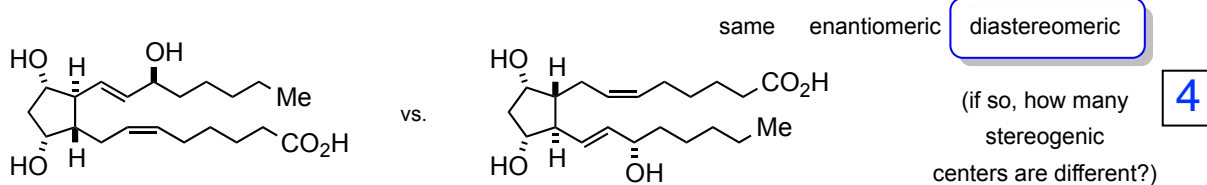
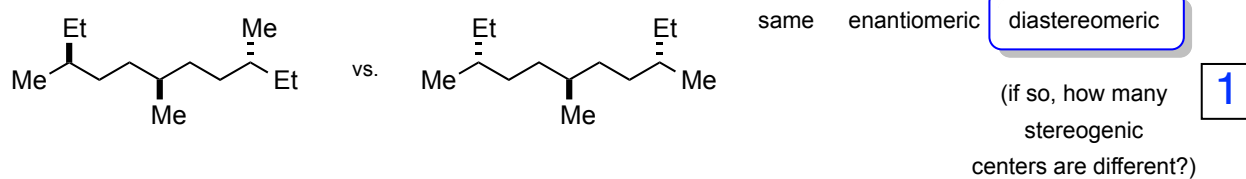
a)			H^+F^-
	ethanol	hydrogen chloride	hydrogen fluoride
b)			
	phenol	4-hydroxypyridine	hydroquinone methyl ether
c)			
	acetic acid	triflic acid	trifluoroacetic acid
d)			
	acetone	nitromethane	ethyl acetate
e)			
	6,6-dimethyl pyran-2,4-dione	dimedone	Meldrum's acid

II. (18 points)

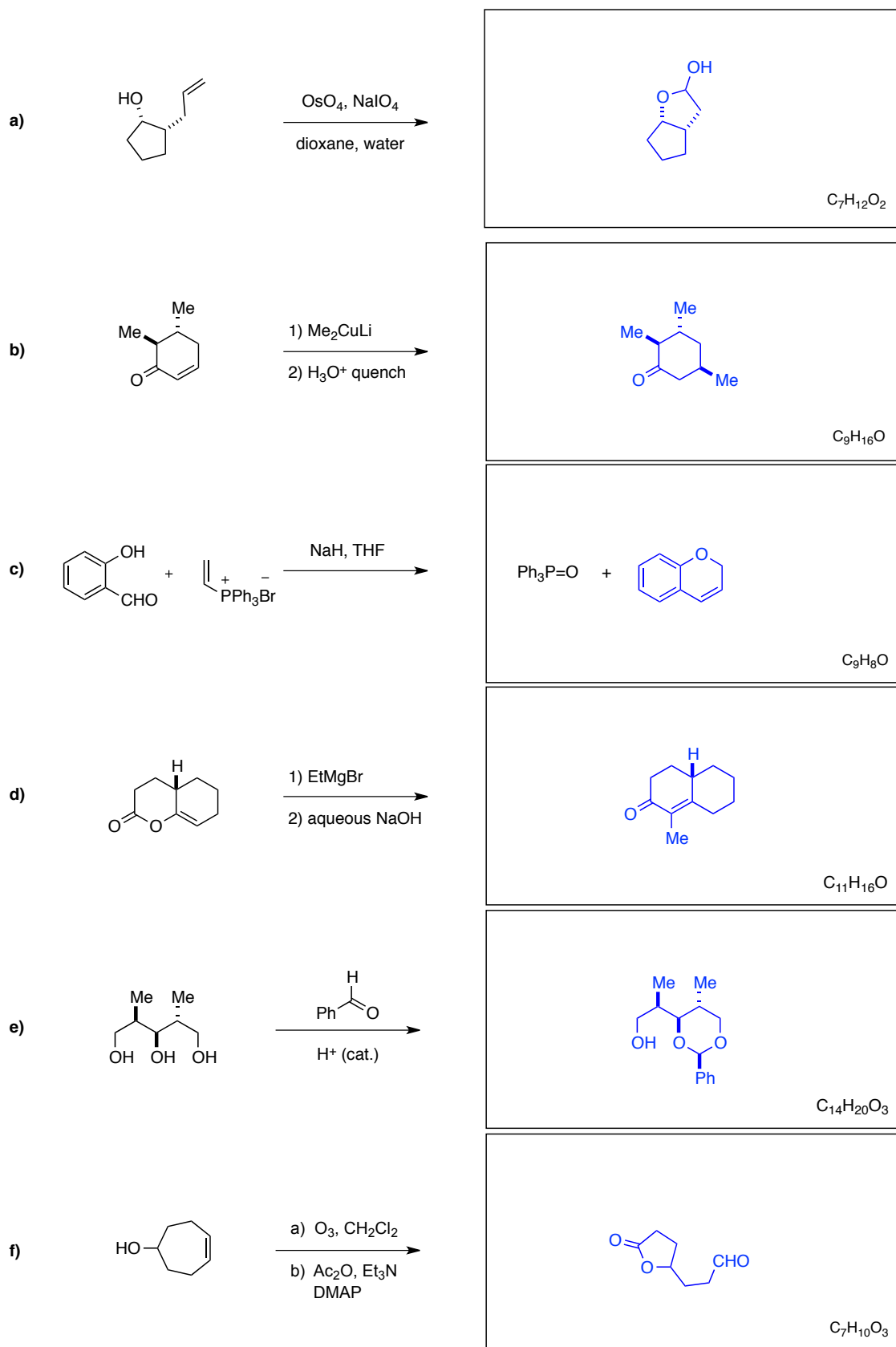
- a) For each pair of similar hydrogen atoms or faces of the sp^2 -hybridized carbon atom, circle the word that properly describes their topicity relative to one another.



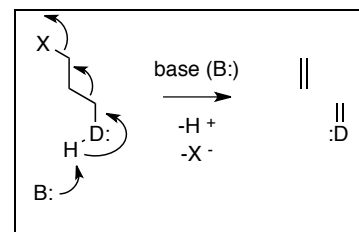
- b) Indicate the relationship (circle the proper word) between each of the following pairs of molecules. For any that are diastereomers, indicate the number of stereogenic centers at which the two have opposite configuration.



III. (24 points) Provide the structure of the major product you expect for each of the following six transformations [a)–f)]. Clearly show the *relative configuration among all stereocenters* for the products of reactions b) and e).

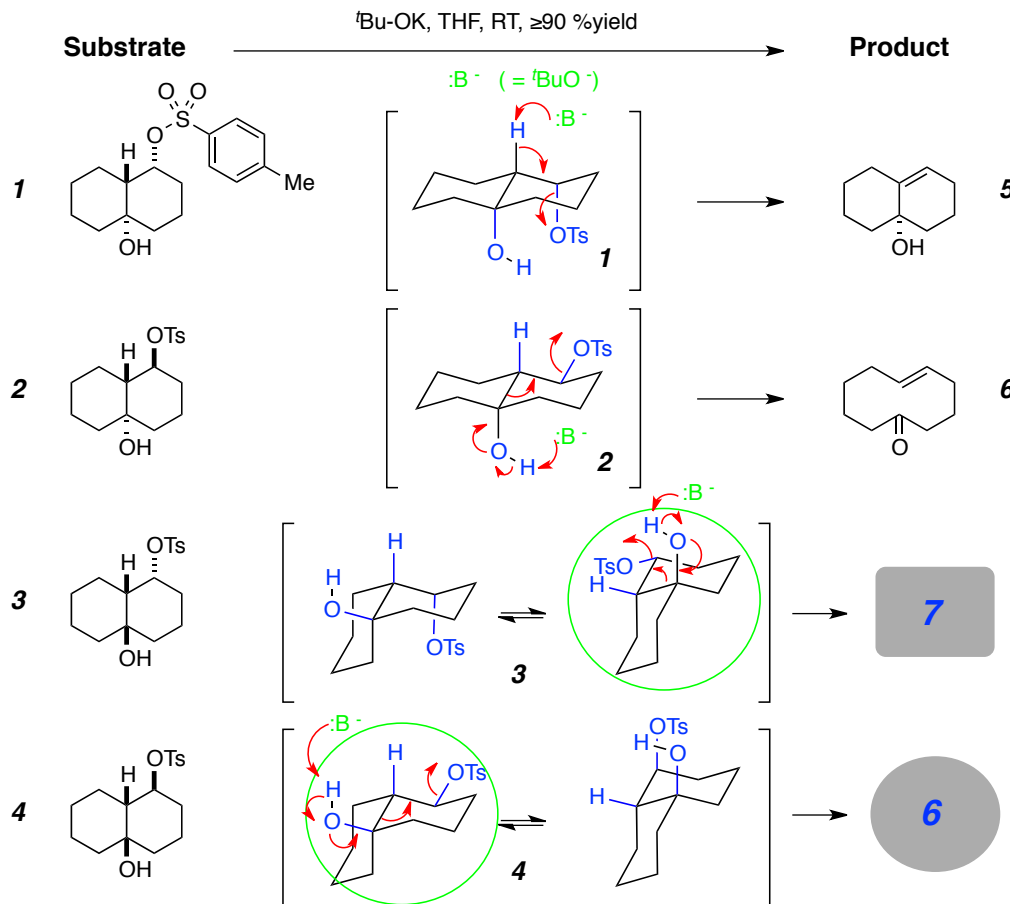
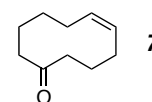


IV. (20 points) This problem is about a reaction known as the Grob fragmentation. The general form of this process is shown in the box to the right, where X is a leaving group and D is a heteroatomic, electron-donating atom (typically, O or N). There is a large stereoelectronic requirement for this fragmentation—the breaking C–C bond needs to be anti to the C–X bond.



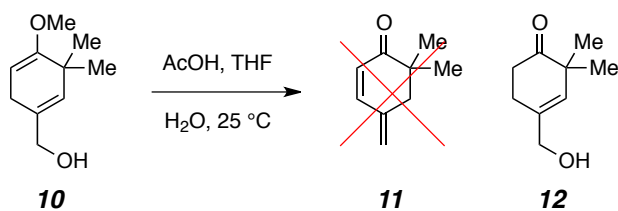
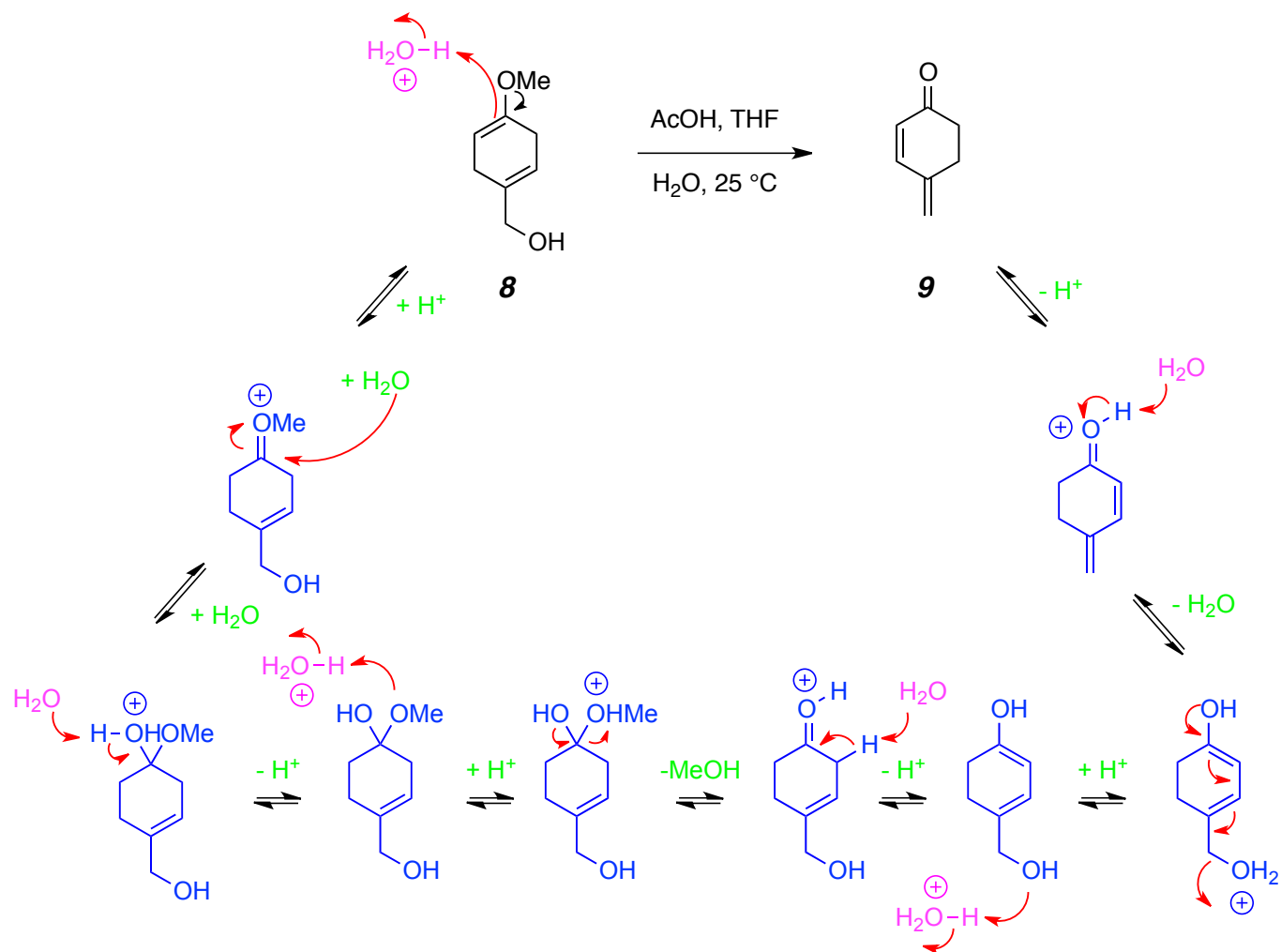
Each of the diastereomeric *p*-toluenesulfonate esters (ROSO₂PhMe ≡ ROTs) **1-4** gives one predominant product (in over 90% yield) when treated with the base, potassium *tert*-butoxide.

- a) Isomer **1** does *not* undergo Grob fragmentation; instead, it gives alkene **5** by a simple E2 elimination process. Add the bridgehead H and OH and the OTs groups in structure **1** to the *trans*-decalin version of that diastereomer [brackets]. Use curly arrows to account for the formation of **5**.
- b) Isomer **2** gives (*E*)-cyclodec-5-enone (**6**), the product of Grob fragmentation. Add the bridgehead H and OH and the OTs groups in structure **2** to the *trans*-decalin version of that diastereomer in [brackets]. Use curly arrows to account for the formation of **6**.
- c) Each of isomers **3** and **4** gives *either 5, 6, or (Z)*-cyclodec-5-enone (**7**) as the product. Add the bridgehead H and OH and the OTs groups in structures **3** and **4** to the *cis*-decalin equilibria, **circle the more stable conformer**. Show the curly arrows for the base-promoted reaction within each of these more stable conformers. Inside the gray box (from **3**) or gray oval (from **4**) write the structure number of the product (*i.e.*, **5, 6, or 7**) that is predicted by each of these analyses.

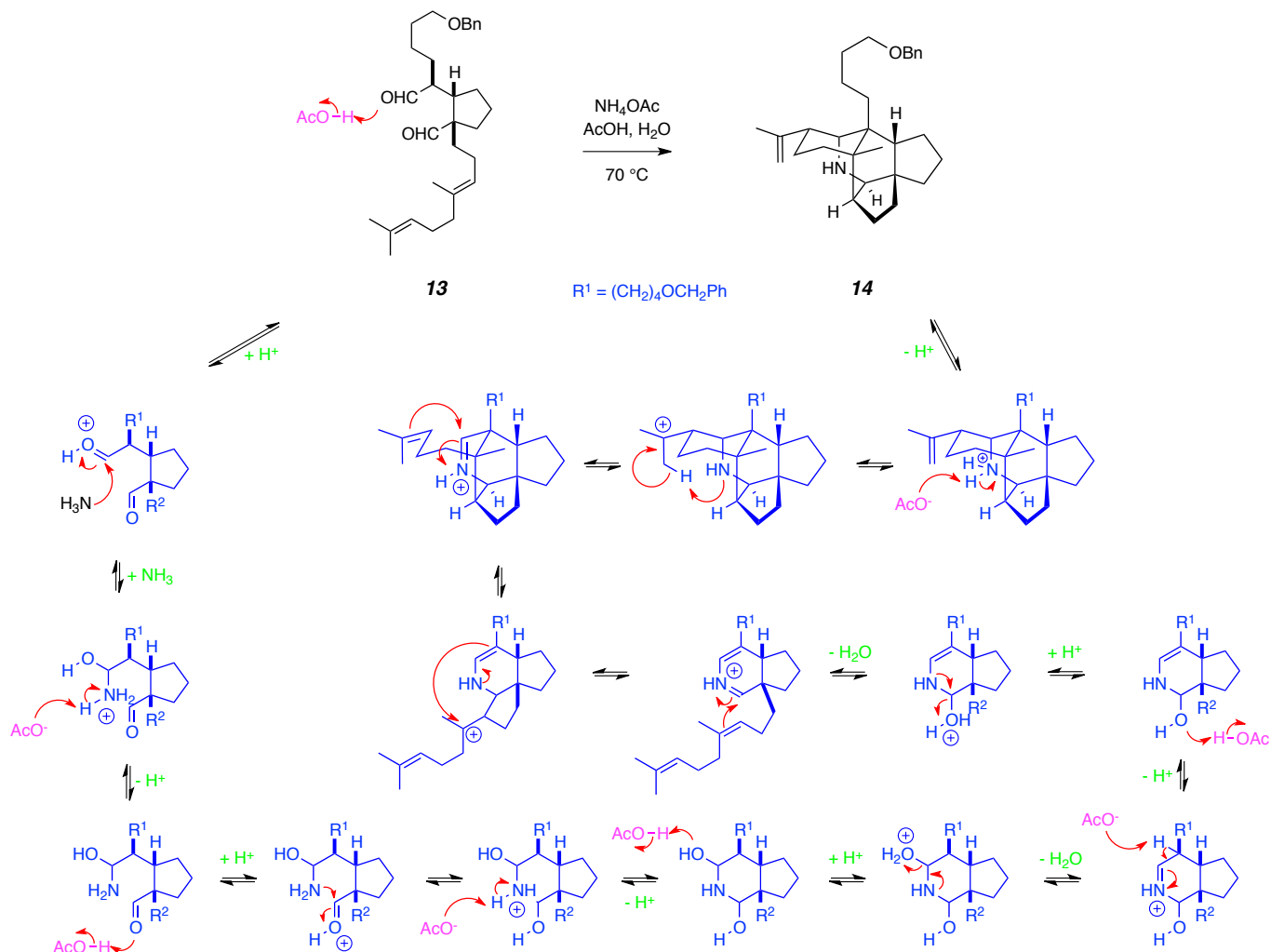


V. (8 pts) Provide a detailed mechanism to account for the conversion of **8** to **9**. Show *ALL* intermediates, equilibria, and bond-making and -breaking steps. There is more than one reasonable pathway that can be envisioned for this acid-catalyzed process; these differ by the order of the fundamental processes involved.

*** For a **5-point bonus**, make sure that the mechanism you provide is consistent with the following two facts: (1) This reaction occurs under mildly acidic (pK_a AcOH = 5) conditions. (2) The *gem*-dimethylated analog **10** does *not* give an appreciable amount of the analogous product **11** under the same conditions but, instead, the ketone **12**.



VI. (20 pts) Based on elegant work from the early 1990s reported from the laboratories of Professor Clayton H. Heathcock, it can be confidently expected that if dialdehyde **13** were treated with ammonium acetate in warm aqueous acetic acid, the polycyclic amine **14** would be formed efficiently. Provide a detailed mechanism to account for this transformation. Show *ALL* intermediates, equilibria, and bond-making and -breaking steps.



If deuterated benzylammonium acetate ($PhCD_2NH_3^+AcO^-$) were to be used instead of ammonium acetate, it can be expected that a deuterated *and reduced* analog of **14**, namely **15**, would be formed instead. Provide a mechanistic rationale (1-2 sentences and 1-2 structures) for this different outcome. [Hint: the deuterated benzaldehyde ($PhCDO$) is the benzyl amine-derived byproduct.]

The benzylic amine **X**, an analog of the penultimate intermediate in the mechanism above, would be formed using $PhCD_2NH_3^+AcO^-$. Intramolecular hydride ion transfer to the tertiary carbenium ion center in **X** would produce the iminium ion **Y**, hydrolysis (or transamination) of which would give **15**.

