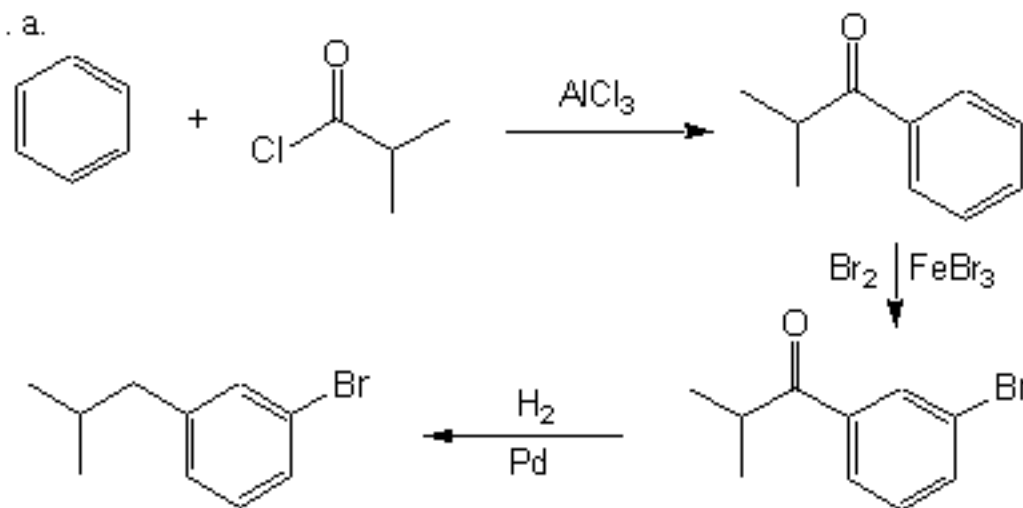


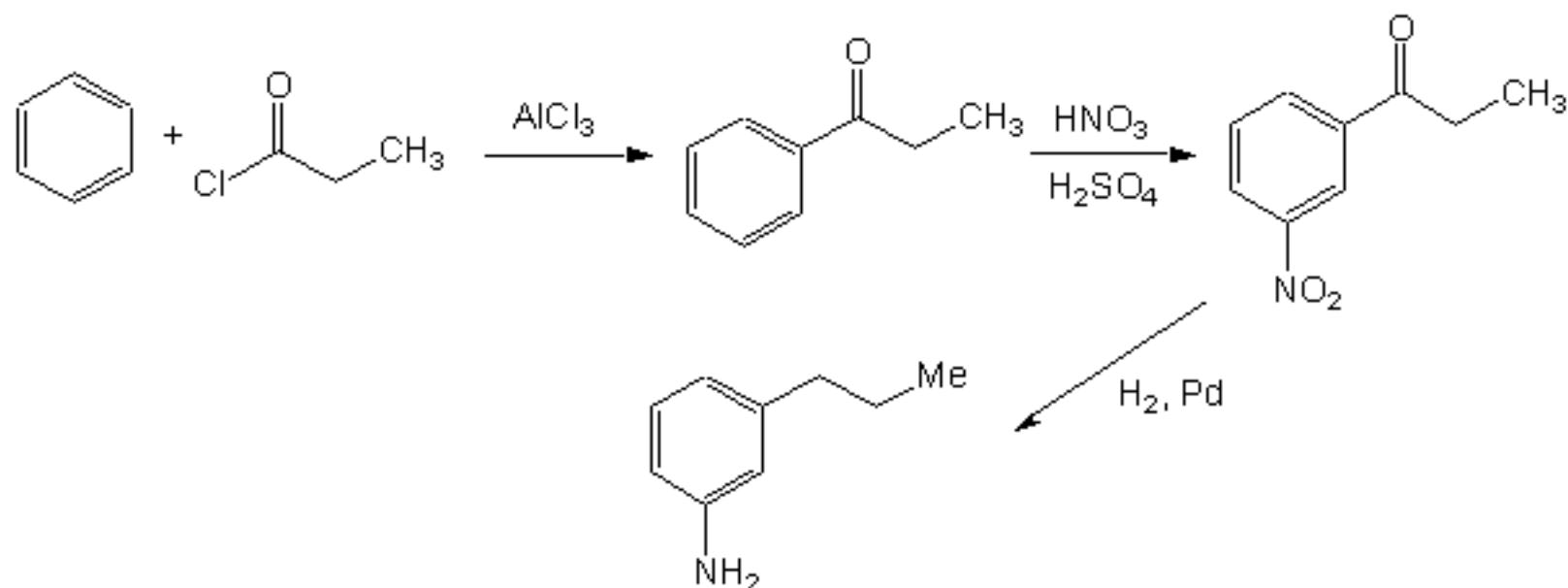
**Organic Chemistry c3444y**  
**Problem Set 2 - Answer Key**

1. a.

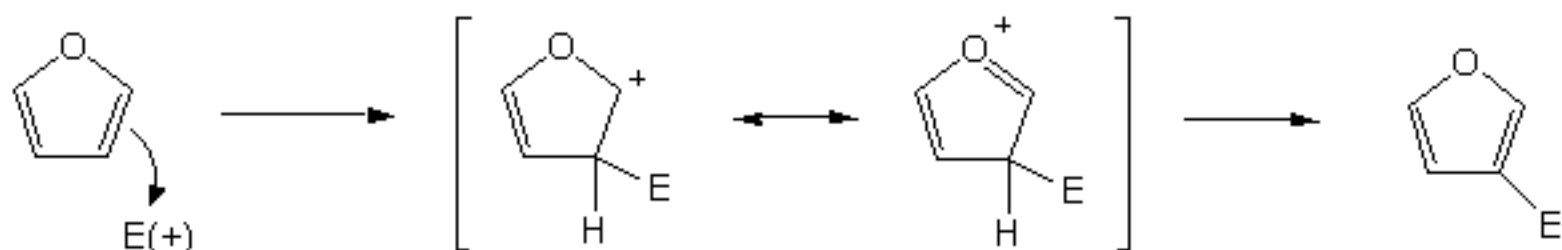
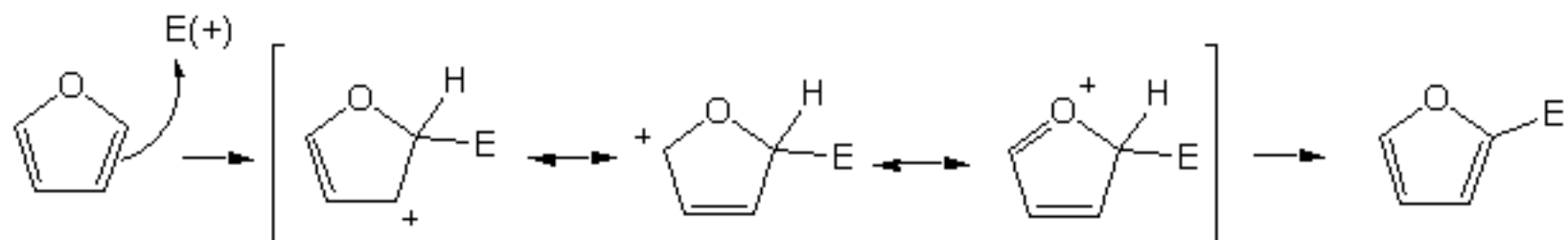
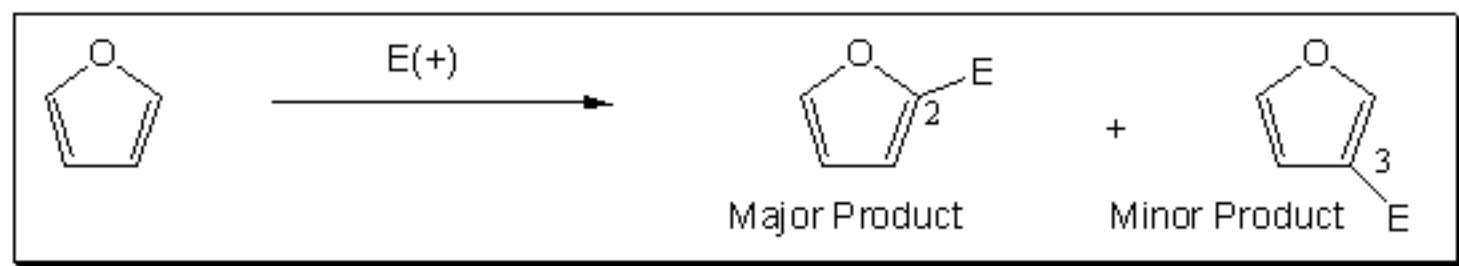


**Analysis of the problem:**  
The target system has two ortho/para directors. There can be no two step synthesis, therefore. We also cannot put an iso-butyl group on directly. So we need a meta director that can be turned into an iso-butyl group.

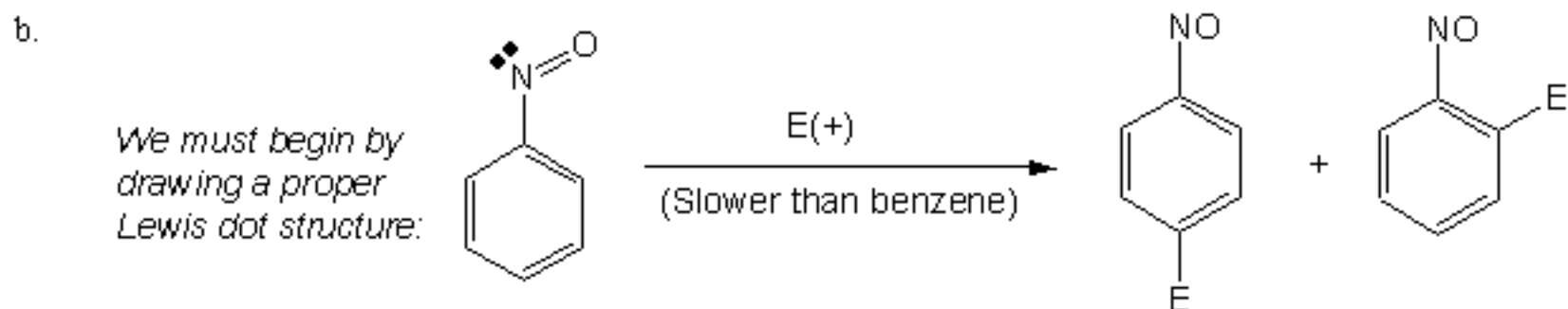
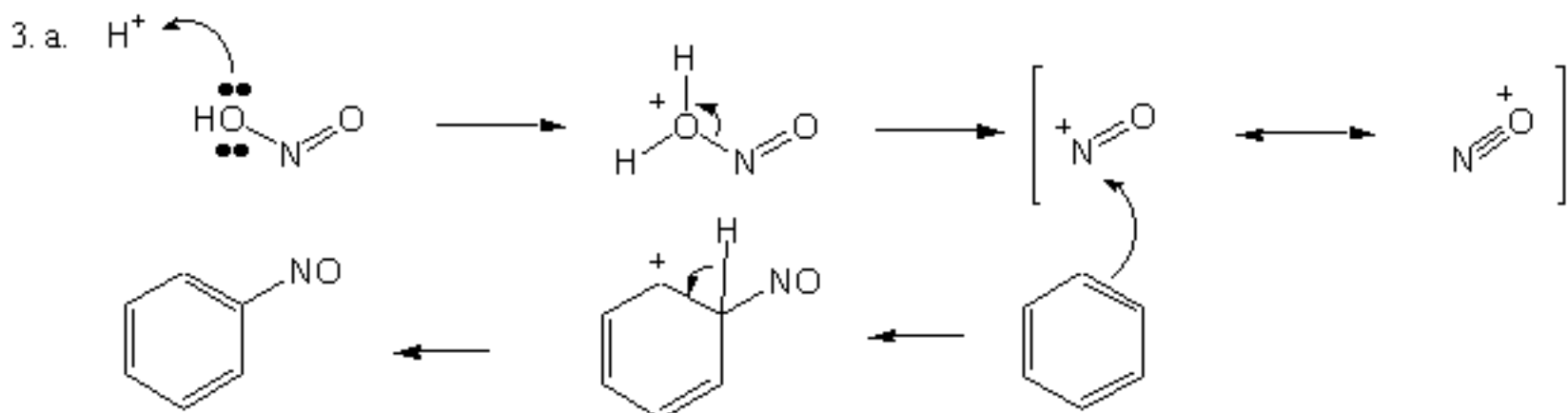
b.



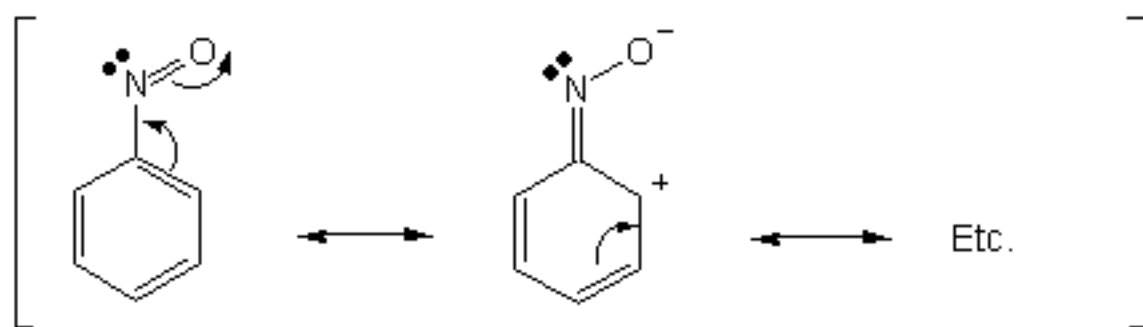
2.



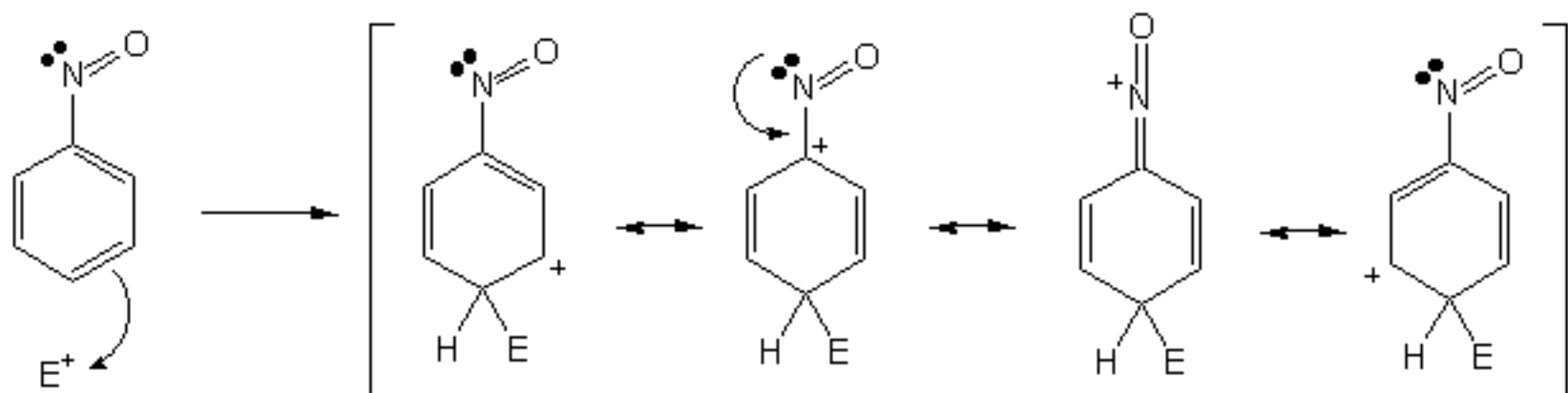
Only two resonance structures here for the intermediate cation. Less delocalization.



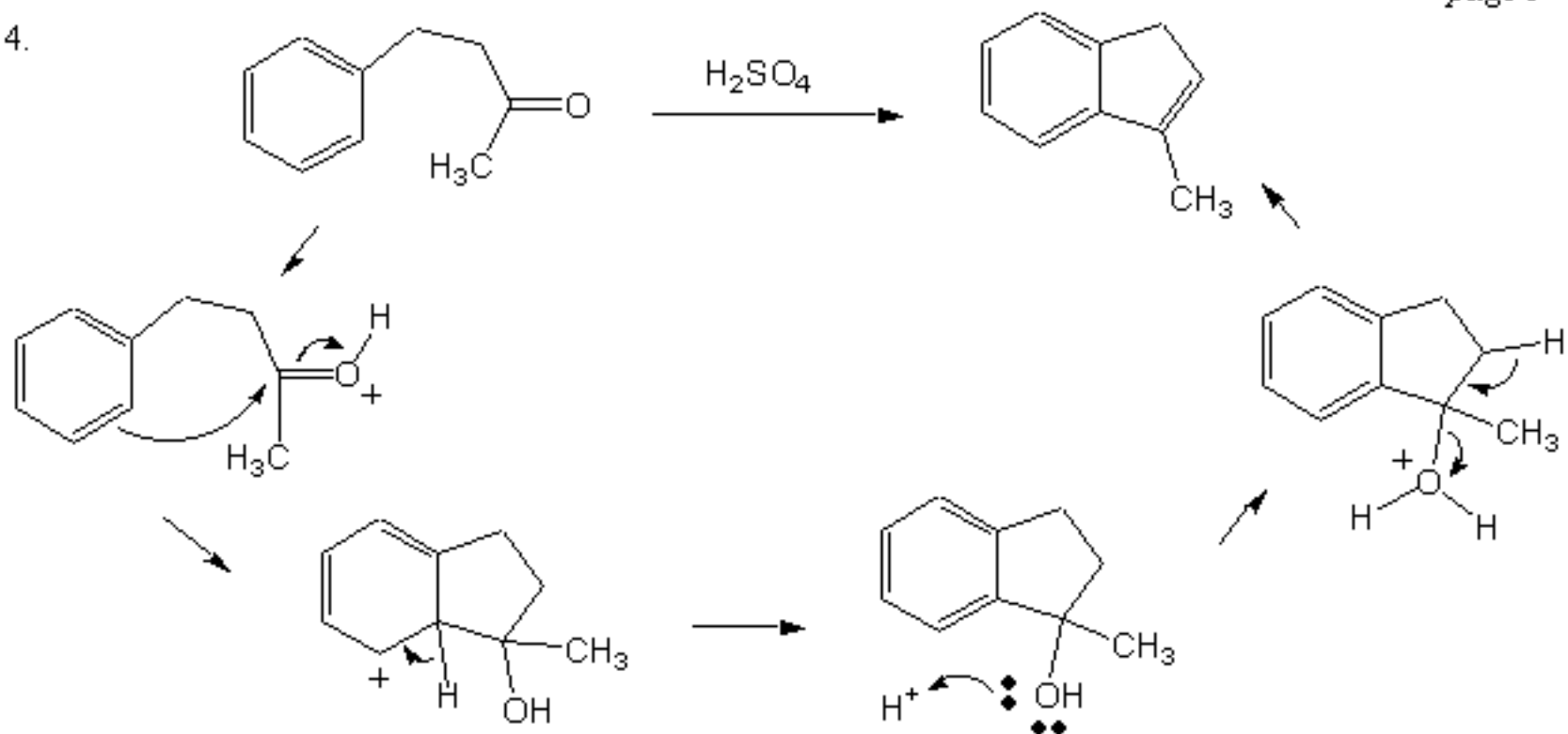
*The NO group is a deactivator because it is an electron withdrawing group, just like a carbonyl or a nitro, due to the electronegative oxygen atom:*



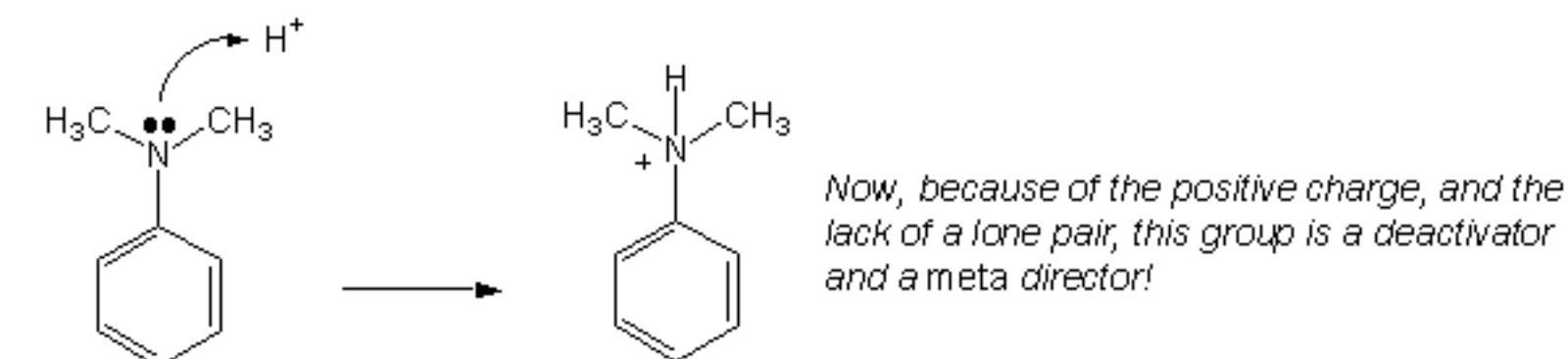
*Like the halogens, however, if it is forced to react it will choose ortho/para to take advantage of the extra resonance structure that we can draw using the lone pair on the N:*



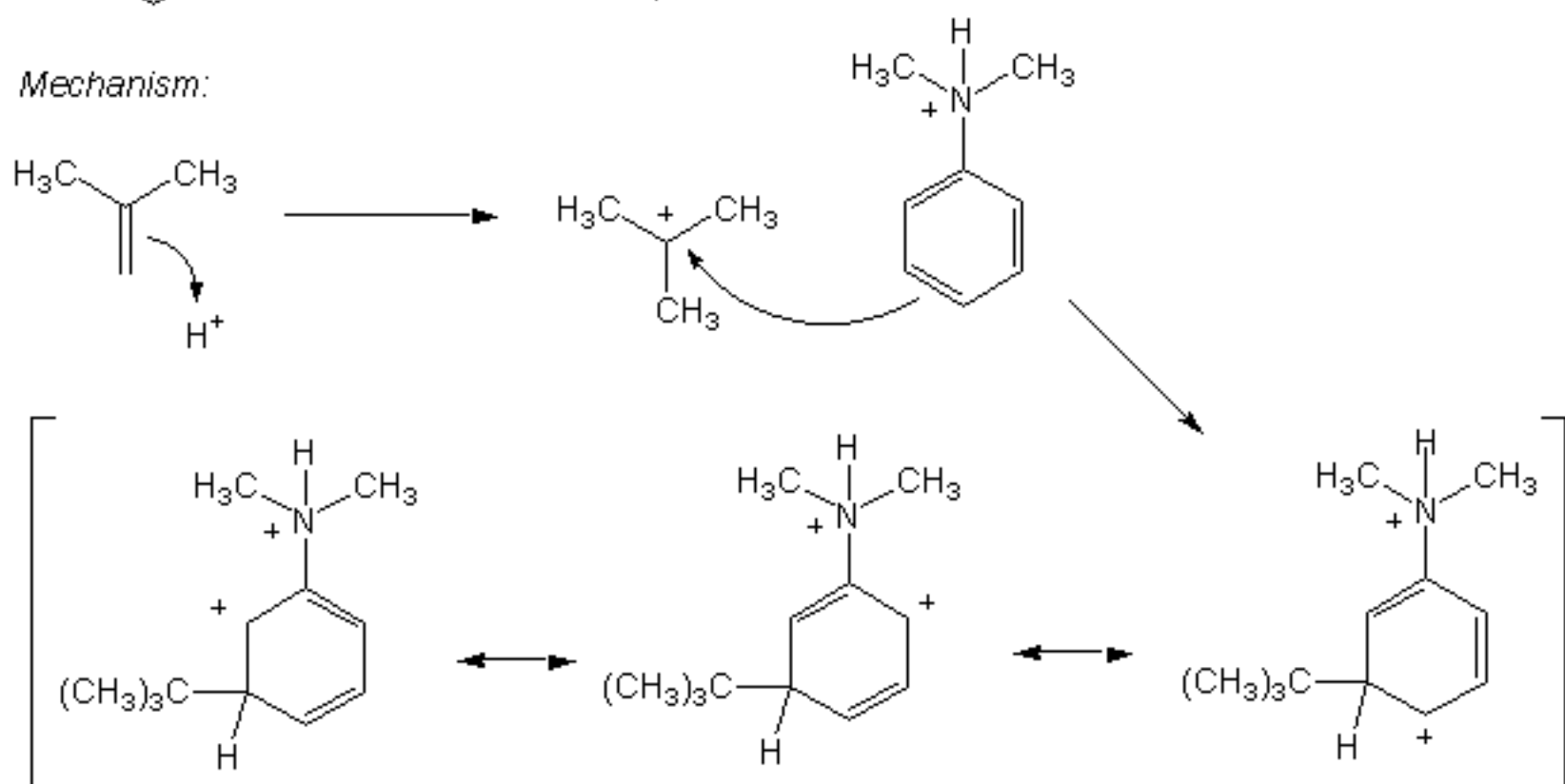
4.



5. The nitrogen of *N,N*-dimethylaniline is a basic nitrogen! Yes, the nitrogen is  $sp^2$  and the lone pair is involved in resonance, but it is not directly involved in the aromaticity. Therefore, it is still basic, albeit weakly. But sulfuric acid is a very strong acid, so what will be the very first thing that will happen:



Mechanism:



Nothing special here, but if it went *para* (or *ortho*), one of the resonance structures would have 2 + charges on neighboring atoms. Thus, the arenium ion formed from meta attack is more stable than that derived from *para* (ortho) attack.