

Ester Synthesis Lab Report

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Introduction

In this lab, we experimented with the production of esters, which is accomplished by refluxing the right alcohol with a parent carboxylic acid while using an acid catalyst. In an ester, an aryl or alkyl group is joined by a carbon to oxygen double bond that is connected to a second oxygen atom. Esters' structural versatility makes them useful in the pharmaceutical business, but they are also widely recognized for carrying the scent of organic materials, making them ideal for perfume, cologne, and even artificial food flavoring.

Results

We continuously poured solution into a container until the chemical diluted and formed a layer. Once the layer had formed, we removed the bottom layer and blended it with other salts. We just kept adding it to a container while creating the top layer. We utilized CaCl_2 pallets that didn't absorb the liquid after removing the layers. Several of our test tubes dried out, but the result—a fragrant smell—remained. (The results are in the image).

starting weight
13.17g

	volume of acetic anhydride (mL)	
acetate	2.3	13.17g
acetate	1.2	
acetate	1.9	14.22
acetate	2.2	
acetate	3.1	13.24
acetate	1.5	
acetate	3.2	

- floral
- Tangerine
- Apple (Sweet Fruit)
- Bananas (Fruit)
- Sweet

Acetic anhydride indicated in the above table in a large dry test tube. Add 3 drops of
alcohol and mix thoroughly. Obtain 2.0 mL of one of the alcohols in a dry small test

tube. Add the alcohol to the acetic anhydride in cold water (to cool the reaction) and add the water with
vigorous mixing between additions. Without cooling, the strongly exothermic
reaction occurs. Then place the tube in hot water at about 70°C for approximately 5

minutes. Mix briefly after each drop. The purpose is to hydrolyze the ester
remaining after the esterification is complete.

4. Add a few drops of a half-saturated sodium chloride solution (3 mL saturated NaCl plus
3 mL water) to the tube and mix thoroughly and vigorously. Then place the tube aside until upper
and lower layers separate. If water alone is used, the two layers would
not separate cleanly. The ester, which is a water-insoluble volatile liquid. In the propyl
acetate test, the two layers separate cleanly with a few minutes. The octyl
acetate test should be run for five minutes to separate the two layers
even after separation.

Questions

1. The SN1 mechanism varies from the SN2 mechanism in several ways. The reactions of primary (and methyl) halides are the slowest, whereas those of tertiary alkyl halides are the quickest. Because the rate law is unimolecular, it only depends on the concentration of the substrate and not the nucleophile (alkyl halide). The most straightforward explanation for the mechanism of this reaction is that it begins with the (rate-determining) loss of a leaving group to generate a carbocation, which is then capable of being attacked by a weak nucleophile at either face, resulting in the loss of stereochemistry.

Sometimes, the SN1 reaction is accompanied by carbocation rearrangements.

2.

3. Peaks are located at 2986, 1745, and 1250 nm for the sp³ CH stretch, C=O stretch, and C-O stretch, respectively. As a result, the molecule possesses a useful ester group.

4.