

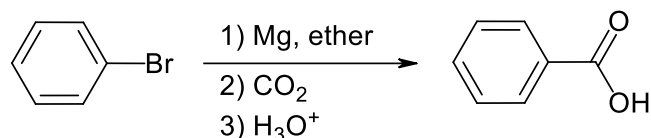
ORGANIC SYNTHESIS: BENZOIC ACID VIA A GRIGNARD REACTION

TECHNIQUES REQUIRED : [Reflux with addition apparatus](#), [rotary evaporation](#)

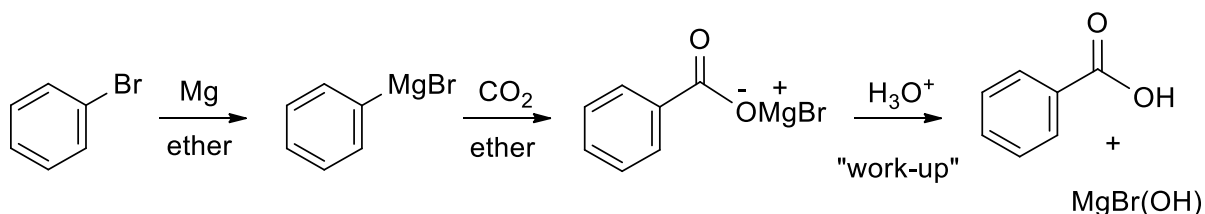
OTHER DOCUMENTS [Experimental procedure](#), [product spectra](#)

INTRODUCTION

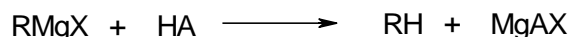
In this experiment you will synthesise benzoic acid using bromobenzene to prepare a Grignard reagent, which is then reacted with solid carbon dioxide, worked-up and purified to give the carboxylic acid. This sequence serves to illustrate some important concepts of practical synthetic organic chemistry : preparing and working with air and moisture sensitive reagents, the "work-up", extractions, apparatus set-up, *etc.* The synthesis utilises one of the most important type of reagents discussed in introductory organic chemistry, organometallic reagents, especially Grignard reagents.



In this reaction, the **Grignard reagent** (an organomagnesium compound), phenylmagnesium bromide is prepared by reaction of bromobenzene with magnesium metal in diethyl ether (the solvent). The Grignard reagent will then be converted to benzoic acid via the reaction of the Grignard reagent with excess dry ice (solid CO₂) followed by a dilute aqueous acid "work-up":



The aryl (or alkyl) group of the Grignard reagent behaves as if it has the characteristics of a carbanion so it is a source of **nucleophilic** carbon which can allow us to make new C-C bonds. It is reasonable to represent the structure of the Grignard reagent as a partly ionic compound, $\overset{\delta^-}{\text{R}} \cdots \overset{\delta^+}{\text{MgX}}$. This partially-bonded carbanion is a very strong base and will react with acids (HA) to give an alkane:

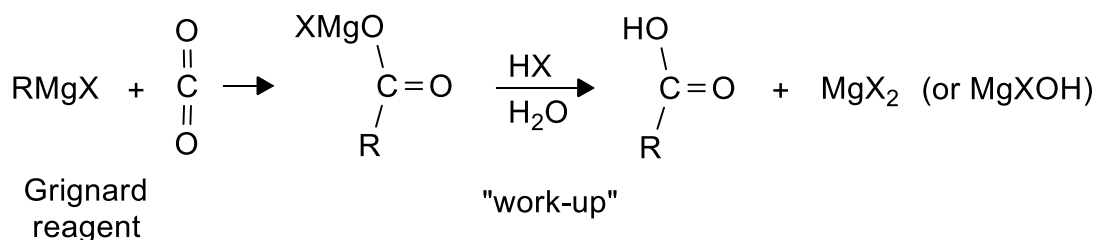


Any compound with suitably acidic hydrogens will readily donate a proton to destroy the reagent. Water, alcohols, terminal acetylenes, phenols and carboxylic acids are just some of the functional groups that are sufficiently acidic to bring about this reaction which is usually an unwanted side reaction that destroys

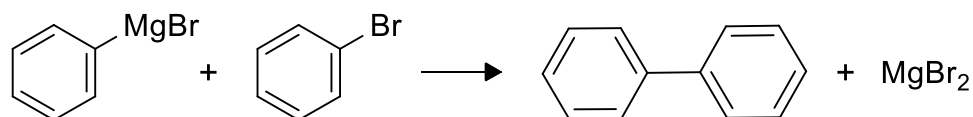
GRIG.2

the Grignard reagent.

The Grignard reagents typically function as a good nucleophiles. A common application has the Grignard reagent adding to the electrophilic carbon of a "carbonyl group" - hence the reactions of Grignard reactions with aldehydes, ketones and certain carboxylic acid derivatives (especially esters) are very important reactions. Here, we can view carbon dioxide as two "back-to-back" carbonyl groups. The Grignard reagent reacts with the CO₂ to form an intermediate carboxylate salt which is not usually isolated. Instead, the intermediate carboxylate salt is neutralised and converted into the product during the "work-up" steps by addition of dilute aqueous acid to create the desired organic compound, the carboxylic acid.



In general, the "work-up" steps are simple acid or base neutralisations that are required in order to convert the initial polar (ionic) intermediate obtained from the reaction into the neutral organic product in a reasonably pure state. The "work-up" steps often neutralise any excess reagents and add acid or base as appropriate (e.g. a reaction that is carried out under basic conditions is usually neutralised with acid) to create the neutral product. Extraction steps may also be used to remove impurities (such as unreacted starting materials) or other by-products. The chief impurities in this Grignard reaction mixture include the hydrocarbon, biphenyl, formed by an unwanted coupling reaction:



This hydrocarbon by-product is easily removed from the main product, benzoic acid, due to the polarity differences and hence solubility differences as the hydrocarbon remains in the organic solvent ether during the extraction steps of the procedure.

REFERENCES

1. <https://www.chem.ucalgary.ca/courses/351/Carey5th/Ch14/ch14-2-2.html>
<https://www.chem.ucalgary.ca/courses/351/Carey5th/Ch19/ch19-2-1.html>