Use of triphenylphosphine-bromotrichloromethane (PPh$_3$-BrCCl$_3$) in the preparation of acylhydrazines, N-methylamides, anilides and N-arylmaleimides from carboxylic acids

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Abstract: In certain countries, many of the reagents used to transform carboxylic acids to acyl halides such as phosphorus trichloride, phosphorus tribromide, phosphorus pentachloride, phosphoryl chloride, thionyl chloride and sulfuryl chloride are difficult to come by. Against this background, the authors developed the reaction system triphenylphosphine (PPh$_3$) – bromotrichloromethane (BrCCl$_3$) to prepare acyl halides in situ. In the following, the use of this reagent combination is joined with the reaction of the in situ prepared acyl halides with nitrogen nucleophiles, specifically with hydrazines, methylamine and anilines. The reaction is also used in an intramolecular variant by the reaction of maleanilic acids to N-arylmaleimides.

Keywords: Appel-type reaction; N-arylmaleimides, acylhydrazines, anilides

1. Introduction

Although there are a number of ways to transform carboxylic acids to acyl halides such as phosphorus trichloride, phosphorus tribromide, phosphorus pentachloride, phosphoryl chloride, thionyl chloride and sulfuryl chloride are not always available in certain countries. Against this background, the authors developed the reaction system triphenylphosphine (PPh$_3$) – bromotrichloromethane (BrCCl$_3$) to prepare acyl halides in situ [1,2]. This reaction system is a variation of the Appel reagent [3], where tetrachloromethane (CCl$_4$) is replaced by the environmentally less hazardous BrCCl$_3$. It was noted that the in situ prepared acyl halides could be reacted with alcohols, and primary and secondary amines to esters and amides, respectively [4]. In the following, the treatment of carboxylic acids is followed by their reaction with hydrazines, methylamine and anilines. Also, further examples [5] will be given to transform maleanilic acids to N-arylmaleimides with the action of PPh$_3$-BrCCl$_3$ in the presence of triethylamine as base.

2. Materials and Methods

3.1. General remarks

Melting points were measured on a Stuart SMP 10 melting point apparatus and are uncorrected. Infrared spectra were measured with a Thermo/Nicolet Nexus 470 FT-IR ESP spectrometer and a Perkin Elmer Spectrum Two spectrometer. $^1$H and $^{13}$C NMR spectra were recorded with a Varian 400 NMR spectrometer (1H at 395.7 MHz, $^{13}$C at 100.5 MHz). The assignments of the carbon signals were aided by DEPT 90 and DEPT 135 experiments (DEPT = Distortionless Enhancement by Polarisation Transfer). The chemical shifts are relative to TMS (solvent CDCl$_3$, unless otherwise noted). Mass
spectra were measured with a JMS-01-SG-2 spectrometer, and with an Agilent QTOF 6540 UHD. Column chromatography, where necessary, was performed on recycled silica gel (S, 0.063 mm – 0.1 mm, Riedel de Haen and Merck grade 9385).

3.2. Starting materials

Benzoic acid (1a), 4-nitrobenzoic acid (1g), 2,4-dichlorophenylacetic acid (6c) (Wako), methylammonium chloride (Merck-Schuchardt), cinnamic acid (4a) (Merck-Schuchardt), 4-methoxycinnamic acid (4b) (Fluka), 4-hydroxybenzoic acid (BDH Chemicals Ltd.), aniline (2d) (Aldrich), p-toluolidin (2b) (4-methylaniline, Merck-Schuchardt), ammonium hydroxide (aq. solution of ammonia, 25 wt%, Sigma Aldrich), and 1-methyl-1-phenylhydrazine (Fluka) were acquired commercially. 3-Chlorobenzoic acid (1b), 4-bromobenzoic acid (1f), 3-bromobenzoic acid (1d), 2-bromobenzoic acid (1c), 2,3-dimethoxybenzoic acid (1h) and 4-methoxybenzoic acid (1e, anisic acid) were prepared from the respective benzaldehydes by oxidation with AgO in basic medium. 4-Alkyloxybenzoic acids were prepared in three steps from 4-hydroxybenzoic acid (transformation to the methyl ester with CH3OH/H2SO4, alkylation of the phenolic function with KOH, Alkyl-I, DMSO [6] and hydrolysis with aq. NaOH in CH3OH). 4-Ethoxy- and 4-propoxyaniline, 18a and 18b, were prepared from 4-hydroxyacetanilide by alkylation to the corresponding 4-alkoxyacetanilides 17a/17b (KOH, Alkyl-I, DMSO [6]) with their subsequent hydrolysis (NaOH, dioxane-MeOH (5:1 v/v) [7]. 4-Triethylamine was distilled over solid KOH and subsequently stored over solid KOH. 2,4-Dinitrophenylhydrazine was dried before use in an oven at 37 °C for 48h.

3.2. General procedures

N-Methyl 3-chlorobenzamide (9a). – To a suspension of triphenylphosphine (PPh3, 980 mg, 3.74 mol) in dry CH2Cl2 (12 mL), BrCCl3 (760 mg, 3.83 mmol) is added dropwise. The solution which turns dark yellow is stirred at rt for 20 min. Then, 3-chlorobenzoic acid (1b, 485 mg, 3.10 mmol) is added, and the resulting mixture is heated at reflux for 25 min. Thereafter, methylammonium chloride (8, 360 mg, 5.41 mmol) is added, and subsequently dry triethylamine (940 mg, 9.30 mmol). The mixture is stirred at reflux for 5h. Thereafter, the cooled reaction mixture is subjected directly to column chromatography on silica gel (CH2Cl2-ether: 10:1) to give 9a as a colorless solid (436 mg, 83%) (mp. 68 °C; Lit. 69-70 °C); IR (KBr/cm−1) v 3338, 3300, 3065, 1638, 1553, 1471, 1407, 1325, 743, 734, 678; 1H NMR (400 MHz, DMSO-d6) δ 2.75 (3H, s, NCH3), 7.46 (1H, dd, J1 = 6.8 Hz, J2 = 6.4 Hz), 7.54 (1H, d, J = 6.8 Hz), 7.74 (1H, d, J = 6.4 Hz), 7.81 (1H, s), 8.58 (1H, bs, NH); 13C NMR (100.5 MHz, DMSO-d6) δ 26.8 (NCH3), 126.2 (CH), 127.3 (CH), 130.8 (CH), 131.4 (CH), 133.6 (C-quat), 136.8 (C-quat), 165.8 (C-quat, CO).

4-Nitrobenzoic acid N-methyl-N-phenyl hydrazide (13d). - To a suspension of triphenylphosphine (PPh3, 980 mg, 3.74 mol) in dry CH2CN (12 mL), BrCCl3 (760 mg, 3.83 mmol) is added dropwise. The solution which turns dark yellow-brown is stirred at rt for 20 min. Then, 4-nitrobenzoic acid (1g, 525 mg, 3.14 mmol) is added and the mixture is stirred at 53 °C for 25 min. Thereafter, N-methyl-N-phenylhydrazine (12, 732 mg, 6.0 mmol) is added and the solution is stirred at 53 °C for 12h. The cooled mixture is concentrated to dryness in vacuo, and the residue is subjected to column chromatography on silica gel (CH2Cl2-ether 10:1) to give 13d (665 mg, 78%) as a light yellow solid; IR (KBr/cm−1) v 3233, 1659, 1597, 1521, 1498, 1345, 1281, 850, 748, 691; 1H NMR (400 MHz, DMSO-d6) δ 3.17 (3H, s, CH3), 6.74 (1H, t, J = 8.0 Hz), 6.81 (2H, d, J = 8.0 Hz), 7.20 (2H, dd, J1 = 8.0 Hz, J2 = 8.0 Hz), 8.09 (2H, d, J = 8.8 Hz), 8.32 (2H, d, J = 8.8 Hz), 10.93 (1H, s, NHCO); 13C NMR (100.5 MHz, DMSO-d6) δ 40.5 (NCH3), 112.9 (2C, CH), 119.1 (CH), 124.2 (2C, CH), 129.3(5) (2C, CH), 129.4 (2C, CH), 139.0 (C), 149.8 (C-quat), 150.0 (C-quat), 164.7 (C-quat, NCO).

N-4-Ethoxyphenylmaleimide (21a). - A solution of triphenylphosphine (1.89 g, 7.21 mmol) and bromotrichloromethane (BrCCl3, 1.56 g, 7.86 mmol) in dry CH2CN (20 mL) is stirred at rt for 30 min., during which in turns dark yellow. Then, maleanilic acid (20a) (1.35 g, 5.76 mmol) is added, and the
The reactions were also carried out successfully with certain oromethane by equilibration with added triethylamine was phase hydrazinium sulfate. For the reaction of the that seen to work well, providing the corresponding N-triphenylphosphine and bromotrichloromethane (BrCCl3). The release of methylamine cinnamic acids reflux. It was noted that the reactions are exothermic. The products were isolated by column to give the anilides imidation was looked at to transform maleianilic acids to hydrazinium salts. The reaction of carboxylic acids with hydrazines. Here, the use of ammonium salts such as methylammonium chloride and the amidation of carboxylic acids conditions the transfer of the bromo transformation of alkanols to haloalkanes with BrCCl3 and the amidation of carboxylic acids similar to those of CCl3 previously, it had been established that the reactions of the modified Appel reagent BrCCl3 were faster than their hydrolysis (Tables 5 and 6). Previously, it had been established that the reactions of the modified Appel reagent BrCCl3-PPh3 are similar to those of CCl3-PPh3. This includes the dehydration of aldoximes and amides to nitriles (1) and the amidation of carboxylic acids (4). Only the actual Appel reaction itself, i.e., the transformation of alkanols to haloalkanes with BrCCl3-PPh3 is more complex, where under certain conditions the transfer of the bromo-substituent vs. the chloro-substituent is not selective (2). In the following, the authors tried to examine whether amidation reactions of carboxylic acids with the less nucleophilic anilines would proceed equally well as with amines. Also, the authors investigated the reactions with hydrazines. Here, the use of ammonium salts such as methylammonium chloride and hydrazinium salts such as hydrazinium sulfate was probed as well as the reaction of carboxylic acids with aq. ammonia in the presence of BrCCl3-PPh3. Finally, the scope of an intramolecular amidation was looked at to transform maleianilic acids to N-arylmaleimides (5).

The reaction of benzoic acids 1, treated with BrCCl3-PPh3 with anilines 2 proceeded straightforward to give the anilides 3 (Table 1). The anilines were added slowly while the reaction mixture was at reflux. It was noted that the reactions are exothermic. The products were isolated by column chromatographic separation on silica gel. The reactions were also carried out successfully with cinnamic acids 4 (Table 2) and phenylacetic acids 6 (Table 3) as substrates.

The release of methylamine in situ, within a reaction mixture of initially carboxylic acid, triphenylphosphine and bromotrichloromethane by equilibration with added triethylamine was seen to work well, providing the corresponding N-methyl amides in acceptable yields (Table 4).

The reaction of aq. ammonia with the in situ prepared acyl halides proceeded equally well, showing that the amidation of the acyl halides proceeded faster than their hydrolysis (Tables 5 and 6).

For the reaction of the in situ prepared acyl halides, three available hydrazines/hydrazine salts available to us were chosen: 1-methyl-1-phenylhydrazine (12), 2,4-dinitrophenylhydrazine (14), and hydrazinium sulfate. 1-Methyl-1-phenylhydrazine is liquid, so its addition to the in situ prepared
acyl halides poses no problem. The 2,4-dinitrophenylhydrazine (14) commercially available to us was dampened with water, most likely to lessen the chance of a detonation of the material. Thus, hydrazine 14 was dried in an oven at 37 °C for 48h before use. Hydrazine itself was released in situ by equilibration of hydrazinium sulfate and triethylamine just as in the case of the use of methylammonium chloride (see above). In all cases, the reaction proceeded reasonably well, delivering acyl hydrazides 13 and 15 (Tables 7 and 8), and in the case of hydrazine itself, 1,2-bis(aroyl)hydrazines 16 were obtained (Table 9).

Finally, our work (5) on using the modified Appel reagent PPh₃-BrCCl₃ for a ring closure of maleianilic acids to N-maleimides was expanded slightly. While previously we had utilized anilines available to us commercially in these reactions, here, we prepared the anilines through a hydrolysis of the corresponding acetamides 17. The general published procedure (7) works very well and allows for the construction of a substituted acetamide before releasing the substituted aniline by hydrolysis.

![Reaction of benzoic acids with anilines in the presence of PPh₃-BrCCl₃](image)

**Table 1.** Reaction of benzoic acids with anilines in the presence of PPh₃-BrCCl₃
Table 2. Reaction of cinnamic acids with anilines in the presence of PPh$_3$-BrCCl$_3$. 

![Chemical Structures](image-url)
Table 3. Reaction of phenylacetic acids with anilines in the presence of PPh₃-BrCCl₃
Table 4. Reaction of substituted benzoic acids with methylammonium chloride in the presence of PPh$_3$-BrCCl$_3$
Proceedings 2019, 3, x FOR PEER REVIEW

1.) PPh₃-BrCCl₃  
CH₂Cl₂

2.) aq. 25 w% 
NH₄OH

Table 5. Reaction of carboxylic acids with aq. ammonia in the presence of PPh₃-BrCCl₃

Table 6. Reaction of cinnamic acids with aq. ammonia in the presence of PPh₃-BrCCl₃
Table 7. Reaction of benzoic acids with 1-methyl-1-phenylhydrazine in the presence of PPh$_3$ – BrCCl$_3$. 
Table 8. Reaction of benzoic acids and 4-chlorophenylacetic acid with 2,4-dinitrophenylhydrazine in the presence of PPh₃ – BrCCl₃.
Table 9. Reaction of benzoic acids with in situ released hydrazine in the presence of PPh₃ – BrCCl₃ to yield 1,2-diacylhydrazines 16.

Table 10. Reaction of maleianilic acids 20 to N-arylmaleimides 21.
5. Conclusion

In conclusion, it can be said that acyl halides, prepared \textit{in situ} by the action of \(\text{PPh}_3\cdot\text{BrCCl}_3\) on carboxylic acids can be transformed further in one pot by the reaction with anilines and hydrazines to anilides and hydrazides, respectively. Also, ammonium and hydrazinium salts can be used when trimethylamine is added as base. Furthermore, the reaction of maleianilic acids with \(\text{PPh}_3\cdot\text{BrCCl}_3\) provides N-arylmaleimides.

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