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Synthesis of α -CF₃-proline derivatives by means of a formal (3 + 2)-cyclisation between trifluoropyruvate imines and Michael acceptors†

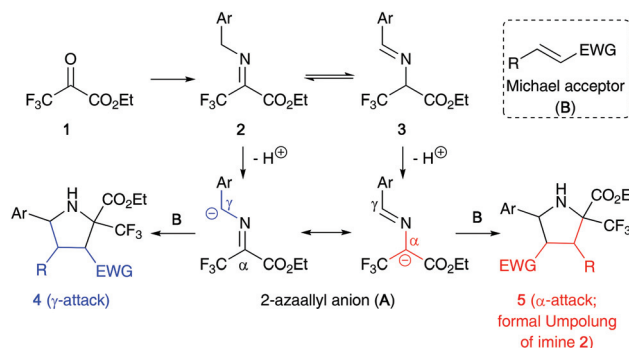
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We herein report the first formal (3 + 2)-cyclisation between 3,3,3-trifluoropyruvate-derived imines and indandione-based Michael acceptors. This reaction gives access to a novel class of spirocyclic α -CF₃- α -proline derivatives with complete control of the diastereoselectivity under phase transfer-catalysed reaction conditions.

Introduction

The synthesis of fluorine-containing α -amino acids became a heavily investigated topic over the last years.^{1–3} The large interest in the hereby accessed fluorinated amino acids is based on the fact that the incorporation of fluorine-containing α - or β -amino acids into peptides and proteins heavily effects their (bio)-chemical and (bio)-physical properties.³ One particularly appealing approach to alter the nature of biologically active molecules is the introduction of a trifluoromethyl group.^{4,5} A broad variety of different synthesis strategies to install either a single fluorine atom, or a polyfluorinated group have been developed in the past.^{6–9} Concerning the synthetic introduction of the trifluoromethyl group, one could either rely on a nucleophilic⁸ or electrophilic⁷ trifluoromethylation of a given compound, or, alternatively, one uses simple CF₃-containing building blocks to access further structural complexity. One commercially readily available CF₃-starting material is ethyl 3,3,3-trifluoropyruvate (**1**), which has been commonly used for further (asymmetric) transformations in the past.^{10–12} Hereby the use of (protected or unprotected) trifluoropyruvate-derived imines is an especially interesting strategy to access (chiral) α -trifluoromethylated α -amino acids.¹¹ Very interestingly, pyruvate **1**-based imines have recently also been used as dipolarophiles for formal (3 + 2)-cyclisations upon reactions with suited 1,3-dipolar compounds.¹² Alternatively, besides their

use as dipolarophiles in (3 + 2)-annulations, imines can also serve as precursors for the *in situ* generation of 1,3-dipolar azomethine ylides or 2-azaallyl anions, which then may react with suited dipolarophiles in a (3 + 2)-fashion.¹³ In this regard also the use of CF₃-containing imines for the construction of 2-CF₃-substituted pyrrolidines has been described.¹⁴ Very interestingly however, and to the best of our knowledge, the trifluoropyruvate-based benzylic imines **2**¹⁵ or the isomeric imines **3** have so far not been used as precursors for the *in situ* generation an utilisation of 1,3-dipolar species. This comes as a surprise, as an (3 + 2)-annulation reaction of 2-azaallyl anions **A** with suited (Michael) acceptors **B** would give access to α -CF₃-containing α -proline derivatives **4** or **5** straightforwardly (Scheme 1). Spurred by the recent progress in the use of *in situ* generated 2-azaallyl anions for (enantioselective) C–C bond forming reactions (often involving a formal Umpolung of the inherent starting imine reactivity),^{16–18} we became interested in investigating the potential of trifluoropyruvate-based benzylic imines **2** for formal (3 + 2)-cyclisations with different Michael acceptors to access a novel class of α -CF₃-substituted



Scheme 1 Targeted use of trifluoropyruvate **1**-based imines to access α -CF₃- α -proline derivatives **4** and/or **5**.

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α -proline derivatives **4** (via γ -attack of the starting imine) or **5** (via Umpolung of the initial imine **2**).

Results and discussion

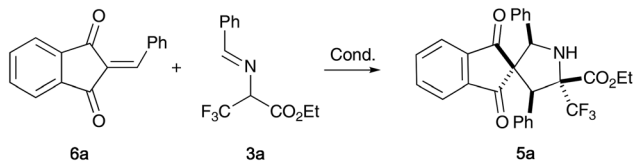
The formation of benzylic imines **2** from pyruvate **1** was investigated in the past, but the synthetic use of these imines has so far only sparingly been evaluated.¹⁹ In these initial reports, Soloshonok and co-workers also found that isolation of imines **2** is hardly possible, as these compounds tend to undergo a very rapid double bond migration to the isomeric benzaldimines **3** under basic and/or higher temperature reaction conditions.¹⁹ Keeping this observation in mind, we reasoned that the α -position of the corresponding 2-azaallyl anions (that will be obtained under basic reaction conditions) will be the preferred nucleophilic site, which should then provide a route towards proline derivatives **5**, rather than **4**.

We started our investigations by carrying out the reaction between the phenyl-based imine **3a** (obtained directly from **1** upon reaction with benzylamine)^{19,20} and the benzyldiene indandione **6a** as the acceptor²¹ (Table 1 gives an overview of the most significant results obtained in this screening). Already our initial experiments showed that the targeted (3 + 2)-annulation is possible, giving the α -attack-derived spiro compound **5a** as the sole cyclization product. It should be noted that in neither case any formation of the corresponding γ -addition product **4** or any classical Michael addition product were observed. As expected, the presence of a simple ammonium salt phase-transfer catalyst (PTC) significantly improved the conversion (compare entries 1 and 2) and the reaction proceeded with full conversion and a promising initial yield and diastereoselectivity in the presence of a catalytic amount of aqueous base (entry 2). In order to improve yield and dr we next screened different bases (in all these cases CH₂Cl₂ was used as the solvent, but we of course also tested other solvents like toluene or THF later on, which however did not allow for better results!).

By increasing the amount of base, we found that solid KOH allows for a slightly higher diastereoselectivity than aqueous KOH (entries 3 and 4). When using different solid bases (entries 4–7) it turned out that LiOH allows for complete control of the diastereoselectivity combined with a reasonable isolated yield of 68% (entry 7). When testing other simple ammonium salt PTCs we found that benzyltriethylammonium bromide performs more or less similarly as tetrahexylammonium bromide (entries 7 and 8) and we later on also realized that the corresponding ammonium chlorides allow for the same outcome. In order to improve the yield further, we increased the amount of base and catalyst (entries 9–11) and finally used a twofold excess of the acceptor **6a** combined with three equivalents of solid LiOH in the presence of 30 mol% ammonium salt catalyst, which gave **5a** in literally quantitative isolated yield on 1 mmol reaction scale (entry 9).

The relative configuration of the major diastereomer was first investigated by NOESY NMR experiments. Here we

Table 1 Identification of the optimum conditions for the diastereoselective (3 + 2)-cyclisation of the azaallyl anion precursor **3a** and acceptor **6a**



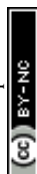
Entry ^a	Cat. ^b	Base	Conv. (%) ^c	Yield ^d (%)	dr ^e
1	—	KOH (50%) (0.1 eq.)	15	n.d.	84 : 0 : 16 : 0
2	THAB	KOH (50%) (0.1 eq.)	>99	57	82 : 12 : 5 : 1
3	THAB	KOH (50%) (1 eq.)	91	74	80 : 12 : 7 : 1
4	THAB	KOH (s) (1 eq.)	93	72	94 : 3 : 3 : 0
5	THAB	CS ₂ CO ₃ (s) (1 eq.)	>99	79	95 : 4 : 1 : 0
6	THAB	K ₃ PO ₄ (s) (1 eq.)	95	51	83 : 10 : 6 : 1
7	THAB	LiOH (s) (1 eq.)	90	68	100 : 0 : 0 : 0
8	TEBAB	LiOH (s) (1 eq.)	>99	70	100 : 0 : 0 : 0
9	TEBAB	LiOH (s) (2 eq.)	>99	70	100 : 0 : 0 : 0
10	TEBAB ^e	LiOH (s) (2 eq.)	>99	75	100 : 0 : 0 : 0
11 ^f	TEBAB ^g	LiOH (s) (3 eq.)	>99	98	100 : 0 : 0 : 0

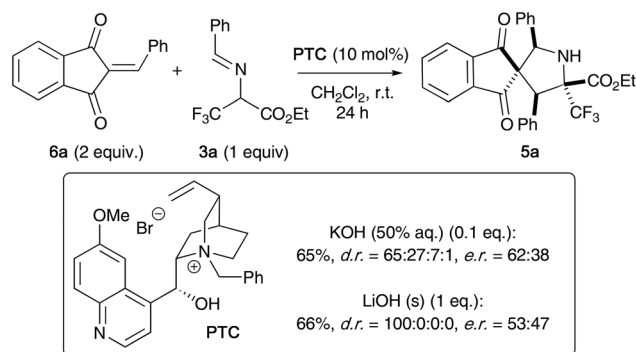
^a Unless otherwise stated all reactions were carried out by stirring a mixture of 0.1 mmol **3a** and 1.2 equiv. **6a** with 10 mol% of catalyst in the presence of base in CH₂Cl₂ at room temperature for 24 hours.

^b THAB (tetrahexylammonium bromide), TEBAB (benzyltriethylammonium bromide). ^c Determined by ¹H and ¹⁹F NMR of the crude product. ^d Isolated yields. ^e Using 20 mol% TEBAB. ^f Using 1 mmol **3a** and 2 equiv. **6a**. ^g Using 30 mol% TEBAB or the analogous chloride TEBAC.

observed NOEs between the two benzylic protons (substantiating a *cis*-configuration of the two phenyl rings) as well as a weak NOE between the ester group and one phenyl group (supporting a *cis* relationship between the ester and the phenyls as well).²⁰ We later on were able to carry out single crystal X-ray analysis of the bromo-derivative **5b**²² (compare with Scheme 3), which unambiguously proved the same relative configuration for that derivative as well, supporting our earlier NOESY results.

Having identified high yielding and highly diastereoselective racemic phase transfer-catalysed conditions for the (3 + 2)-cyclisation between **3a** and **6a**, we next screened a variety of different chiral ammonium salt phase transfer catalysts under different conditions for this reaction.^{20,23} Unfortunately however, all these efforts were rather disappointing, as we were not able to obtain any reasonable enantioselectivity for this reaction. Scheme 2 gives the best result obtained with a Cinchona alkaloid-based chiral phase transfer catalyst, but a variety of other catalyst systems under different conditions were tested as well, but none of them gave any satisfying selectivity (as shown in the online ESI†).²⁰

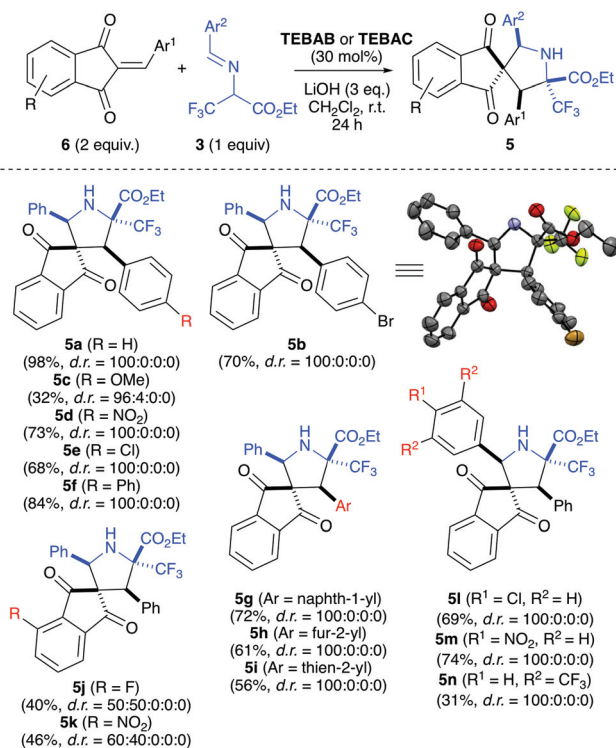




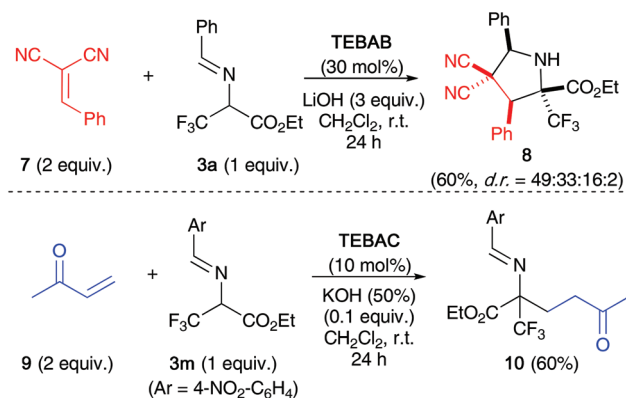
Scheme 2 Attempted chiral phase transfer-catalysed enantioselective (3 + 2)-cyclization.

As we were not successful in developing a satisfyingly enantioselective protocol, we investigated the application scope for the reaction between imines **3** and 1,3-indandiones **6** using achiral ammonium salt PTCs only (Scheme 3).

A variety of acceptors with different Ar¹-groups were tolerated well (giving products **5a–5i**). Only the methoxy-containing **5c** was obtained in a lower yield, mainly because of a significantly slower conversion in that specific case. As mentioned above, **5b** could be analysed by single crystal X-ray analysis and the relative configuration of the other products was assigned in analogy. When using alternative Ar²-groups on the donor side, the products **5l** and **5m** were obtained in similar yields



Scheme 3 Application scope of the diastereoselective (3 + 2)-cyclization of imines **3** and acceptors **6**.



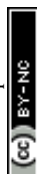
Scheme 4 Reaction of imines **3** with alternative acceptors **7** and **9**.

and with the same high diastereoselectivity. Surprisingly, the bis-CF₃-containing **5n** was obtained in a lower yield only, mainly because of the accompanying formation of unidentified side-products (most probably originating from background hydrolysis of the starting imine). Unfortunately, the introduction of other substituents on the indandione side resulted in slightly reduced yields (see products **5j** and **5k**). Furthermore, these products were also obtained as mixtures of diastereomers (differing in the configuration of the spiro carbon). The fact that the configuration of this additional stereogenic center was not controlled in this otherwise highly diastereoselective reaction came as a surprise, and unfortunately we were not able to overcome this issue.

Finally, we also tested if other Michael acceptors may be used for this (3 + 2)-cyclisation with imines **3**. As shown in Scheme 4, the arylidene malonitrile **7** gave the (3 + 2)-product **8** as well, but with very low diastereoselectivity and in this case also significant amounts of different not identified impurities were formed. In sharp contrast, methyl vinyl ketone **9** was the only acceptor so far that did not give the cyclisation product, but rather underwent a reasonably high yielding Michael addition when reacted with the NO₂-containing imine **3m** (the same conjugated addition reaction, but significantly slower, was observed with the phenyl-based **3a**).

Conclusions

The reaction between 3,3,3-trifluoropyruvate (**1**)-derived imines **3** and indandione-based Michael acceptors **6** allows for a formal (3 + 2)-cyclisation which gives access to an unprecedented class of α-CF₃-α-proline derivatives **5**. By choice of the proper phase transfer-catalysed reaction conditions (*i.e.* the used base) this reaction performs with literally complete diastereoselectivity. Unfortunately, the use of chiral ammonium salt catalysts turned out to be not satisfying so far. Other acceptors were tested as well and arylidene malonitrile **7** undergoes this type of (3 + 2)-cyclization as well, but with relatively poor diastereoselectivity, while methyl vinylketone (**9**) acted as a Michael acceptor only.



Experimental details²⁰

General procedure

The arylidene indandiones **6a–k** (0.2 mmol) were dissolved in dry CH₂Cl₂ (2 mL), followed by the addition of LiOH (0.3 mmol, 7.2 mg), TEBAAC (30 mol%, 0.03 mmol, 7.8 mg) and imines **3a–d** (0.1 mmol). The reaction mixture was stirred for 24 h at room temperature followed by filtration over a pad of silica and washing with Et₂O. After evaporation of the solvent, the product was purified by column chromatography with a gradient of heptanes and EtOAc (20 : 1–10 : 1–5 : 1) to yield products **5a–n** in the reported yields and diastereomeric ratios.

Analytical data for **5a** (this compound was also synthesised on 1 mmol scale)

¹H NMR (300 MHz, CDCl₃, 298 K) δ 7.76–7.73 (m, 1H), 7.60–7.52 (m, 3H), 7.15–7.06 (m, 10H), 4.93–4.83 (m, 2H), 4.65 (s, 1H), 4.31–4.20 (m, 1H), 3.96–3.86 (m, 1H), 0.94 (t, J = 7.1 Hz, 3H) ppm; ¹³C NMR (176 MHz, CDCl₃, 298 K) δ 199.4, 199.4, 166.7, 142.6, 142.1, 135.9, 135.6, 133.5, 133.5, 130.3, 128.4, 128.4, 128.2, 127.9, 126.7, 126.2 (q, J = 298 Hz), 124.1, 122.0, 122.9, 74.6 (q, J = 28 Hz), 72.1, 71.3, 63.0, 60.7, 14.2, 13.4 ppm; ¹⁹F-NMR (282 MHz, CDCl₃, 298 K): δ = –74.22 (s, 3F) ppm; HRMS (ESI): *m/z* calcd for C₂₈H₂₂F₃NO₄ [M + H]⁺ = 494.1574, found: 494.1580.

Conflicts of interest

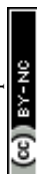
There are no conflicts to declare.

Acknowledgements

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