Article

β -Nitroacrylates: New Key Precursors of Indole-2-Carboxylates via Fischer Indole Synthesis

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Received: 5 November 2019; Accepted: 23 November 2019; Published: 28 November 2019

Abstract: Indole 2-carboxylates are very important scaffolds that are widely investigated for their activities and are used as key intermediates of biologically active molecules. Herein, we report a new procedure for the preparation of this class of derivatives, via Fischer indole synthesis, starting from β -nitroacrylates and arylhydrazines. The protocol permits the production of the title targets in satisfactory overall yields, avoids any wasteful aqueous work-up, and has with evident advantages from a sustainability point of view.

Keywords: Indoles; β -nitroacrylates; Fischer indole synthesis; heterocycles; conjugate addition

1. Introduction

The indolic system is one of the most important and versatile nitrogen containing heterocycles; it is widely spread in nature, such as in food, flowers, in a multitude of biologically active molecules, and is probably represented as one of the “privileged structures” of the main scaffolds for the discovery of new drug candidates [1,2]. For all these reasons, a growing number of synthetic approaches for its preparation and derivatization have been reported in the literature [3–5]. Nowadays, notwithstanding the development of new methodologies concerning the ex-novo preparation of indoles, the historical Fischer indole synthesis (introduced by Emil Fischer in 1883) remains one of the pillars for the direct production of functionalized indoles (Scheme 1) [6,7].

Scheme 1. Fischer indole synthesis.

This reaction is based on the indolization of arylhydrazones under acidic conditions, and over the years, several studies have been done concerning the discovery and use of even-more efficient acid catalysts [8]. However, only few efforts have been directed toward the preparation of arylhydrazones, which are mainly prepared from ketones and arylhydrazines (Scheme 2) [9–11].

Scheme 2. Synthesis of arylhydrazones.

While simple ketones are cheap and readily available, the functionalized ones very often require a multistep synthesis and harsh reaction conditions. Among them, α -ketoesters are of particular interest since they are key precursors of alkyl indole-2-carboxylates, a very important class of indole that is widely investigated for its activities and is exploited as a strategic intermediate of biologically active targets, such as the NNRTI Delavirdine and its analogues [12–17]. Recently, following our study concerning the use of β -nitroacrylates **1** as a useful building block of heterocyclic systems [18–21], we reported a new protocol for synthesizing **5**, starting from **1** and *o*-bromoanilines, based on a palladium catalyzed Heck coupling [22] (Scheme 3, Way b). Now, in continuing this study with the aim of avoiding the use of precious metal (Pd), we found **1** to be a valuable alternative of α -ketoesters for the Fischer indole synthesis (Scheme 3, Way a).

Scheme 3. Synthesis of alkyl indole-2-carboxylates starting from β -nitroacrylates.

2. Materials and Methods

2.1. General Section

OXFORD NMR S400, Varian Mercury Plus 400, Oxford, UK, equipped with workstation Sun Blade 150, software VNMRJ 1.1d, and operating system Solaris 9. ^1H NMR analyses were recorded at 400 MHz and ^{13}C NMR analyses were recorded at 100 MHz. Ir spectra were recorded with a Spectrum Two FT-IR spectrometer, Waltham, MA, USA equipped with ZnSe window, Dynascan Interferometer, detector type LiTaO₃, and Spectrum 10 software. Microwave irradiations were performed by Biotage® Initiator. Microanalyses were performed with a CHNS-O analyzer Model EA 1108 from Fisons Instruments. GS-MS analyses were obtained on an Agilent GC(6850N)/MS(5973N), Stevens Creek Blvd, Santa Clara, CA, USA, EI technique (70 eV), GC/MSD software, and an HP-5MS column, 30m, Id 0.25 μm , film thickness 0.25 μm . Amberlyst 15 was purchased from Sigma–Aldrich and was purified by soaking it in methanol for 24 h and then washing it with fresh methanol and THF and dried under a vacuum.

2.2. Preparation of Starting Materials

β -Nitroacrylates **1** were synthesized by the Henry reaction–elimination process, starting from nitroalkanes and alkyl glyoxalates [23,24]. Arylhydrazines were purchased from Sigma–Aldrich as hydrochloride salts and were converted into the free bases by treatment with potassium carbonate in water [25].

2.3. General Procedure for the Preparation of Compounds **5**

A mixture of β -nitroacrylate **1** (1 mmol) and arylhydrazine **2** (1 mmol) was stirred, under solvent-free and promoter-free conditions, at room temperature for 2 h. Successively, acetonitrile (6 mL) and TMG (2 mmol, 0.230 mg) were added, and the resulting solution was stirred for a further 4 h. Then, the acetonitrile was removed under a vacuum, and for the purposes of removing the TMG, the reaction crude was filtered through a short pad of silica (3 g) and washed with ethyl acetate (25 mL). After evaporation under the vacuum of ethyl acetate, the mixture of tautomers **4**, **4'**, and **4''** were dissolved in 10 mL of the appropriate alcohol, treated with 2 g of Amberlyst 15, and irradiated by the Biotage® Initiator⁺ at 90 °C for 4 h. Finally, the resin was filtered off and it was washed with fresh ethyl acetate (15 mL) and the crude product **5**, which was obtained after removal of the solvent at reduced pressure, and was purified by flash chromatography column (hexane–dichloromethane 6:4).

3. Results and Discussion

This new approach (Scheme 3, Way a) involves two different steps: (i) A one-pot Michael addition/elimination process, in which the arylhydrazines **2** reacts with **1** to form adduct **3**, which under basic conditions, eliminates nitrous acid and provides intermediate **4**; and (ii) an acidic promoted indolization of **4** into the title target **5**.

3.1. Optimization of the 1st Step

In order to optimize the entire protocol, we first investigated the Michael addition/elimination one-pot process using phenylhydrazine **2a** and the β -nitroacrylate **1a** as reagents. Preliminary studies highlighted the almost quantitative addition of **2a** to **1a** under solvent-free and promoter-free conditions at room temperature (2 h). Further investigations (in terms of solvents, bases, and reaction temperatures) were dedicated to nitrous acid elimination, and the best result was obtained using two equivalents of TMG in MeCN at room temperature (4 h, entry f). Instead of obtaining the pure adduct **4a**, a mixture of tautomers **4a**, **4a'**, and **4a''** were isolated in a ratio of approximately 54:13:33. The ratio was calculated by ¹H-NMR on the basis of the characteristic signal of each tautomer, as depicted in Figure 1. Nevertheless, all tautomers were suitable for the Fischer indole synthesis (Table 1).

Figure 1. Characteristic ¹H-NMR signal of tautomers **4a**, **4a'**, and **4a''** (tautomeric ratio 54:13:33).

Table 1. Optimization studies concerning the addition/elimination step.

Reaction scheme: 1a (ethyl 2-ethyl-3-nitroacrylate) reacts with (1 eq.) PhNHNH₂ under Neat conditions for 2 h to form intermediate 3a (ethyl 2-ethyl-3-nitro-2-phenylhydrazine-3-carboxylate) quantitatively. Intermediate 3a then reacts with Base in Solvent at Temp. (°C) to form a mixture of tautomers 4a, 4a', and 4a''.

Entry	Base	Solvent	Temp.	Time (h)	Yield (%) of 4a-a''
A	DBU (1 eq.)	MeCN	rt	2	23
B	TMG (1 eq.)	MeCN	rt	4	55
C	KF/Al ₂ O ₃ (1 eq.)	MeCN	rt	24	— ¹
D	PS-Carbonate (1 eq.)	MeCN	rt	24	— ¹
E	PS-TBD (1 eq.)	MeCN	rt	4	35
F	TMG (2 eq.)	MeCN	rt	4	75
G	TMG (2 eq.)	MeCN	50 °C	1	60
H	TMG (3 eq.)	MeCN	rt	4	71
I	TMG (2 eq.)	THF	rt	4	35
J	TMG (2 eq.)	EtOAc	rt	4	49
K	TMG (2 eq.)	DCM	rt	4	21

¹ No reaction occurs.

3.2. Optimization of the 2nd Step

Successively, we faced off the indolization of tautomers **4a-a''** into the alkyl indole-2-carboxylate **5a**. In this context, we first performed the addition/elimination one-pot process, then the TMG was removed by fast filtration through a short pad of Silica, and finally, after the removal of the solvent under a vacuum, the crude **4a-a''** were subjected to a variety of acidic conditions. Initially, inspired by the literature, we tested PPA, ZnCl₂/AcOH, SOCl₂/EtOH, and *p*-toluenesulfonic acid/EtOH [26–29]. Only the latter system was efficient for the indolization reaction (3eq. *p*-TsOH, EtOH, reflux, 8 h, 32%). Based on this result and following our studies concerning the use of heterogeneous systems, we repeated the reaction using 1g/mmol of Amberlyst 15, a sulfonic acid type based upon a styrene-DVB copolymer [30]. Under these conditions, **5a** was recovered in a similar yield (30%), but with the remarkable advantage of minimizing the work-up step to an easy filtration, thus avoiding the classic tedious and wastefully aqueous work-up. Further tests permitted the optimization of reaction conditions (Table 2), and the best yield of **5a** (over the two steps) was recorded, under microwave irradiation at 90 °C for 4 h, using 2 g/mmol of Amberlyst 15 (Entry k, 54%).

Table 2. Optimization studies concerning the indolization step.

Entry	Amberlyst 15 (g/mmol)	Temp. (°C)	Time (h)	Yield (%) ¹ of 5a
<i>a</i>	1	reflux	8	30
<i>b</i>	1.5	reflux	8	34
<i>c</i>	2	reflux	8	38
<i>d</i>	2.5	reflux	8	39
<i>e</i>	2	60	8	15
<i>f</i>	2	100 ²	8	40
<i>g</i>	2	reflux	16	45
<i>h</i>	2	reflux	24	53
<i>i</i>	2	reflux	32	51
<i>j</i>	2	80 ³	4	42
<i>k</i>	2	90 ³	4	54
<i>l</i>	2	100 ³	4	49

¹ Overall yield of the pure isolated product. ² Reaction performed in a sealed vessel. ³ Reaction performed in a sealed vessel under microwave irradiation.

3.3. Substrate Scope Investigation

Lastly, to achieve the aim of verifying the generality of our protocol, we tested a variety of β -nitroacrylates and arylhydrazines under the optimized reaction conditions (Table 3). In all cases, the products were isolated from satisfactory to good overall yields (38–55%), even in the presence of versatile and reactive functionalities. In particular, it was possible to prepare indoles functionalized on the C-3 chain that were suitable for further manipulations, such as **5e–g** and **5j**, which contained a phenyl ring, a double and triple bond, and an ester group, respectively.

Table 3. Synthesis of indoles **5a–m**.

Entry	R^2	R^3	G	Solvent	Yield (%) ¹ of 5
<i>a</i>	Et	Et	H	EtOH	5a 54
<i>b</i>	$\text{CH}_3(\text{CH}_2)_4$	Et	H	EtOH	5b 52
<i>c</i>	$\text{CH}_3(\text{CH}_2)_8$	Et	H	EtOH	5c 50
<i>d</i>	Et	Et	7-Me	EtOH	5d 55
<i>e</i>	$\text{Ph}(\text{CH}_2)_2$	Et	H	EtOH	5e 43
<i>f</i>	$\text{CH}_2=\text{CH}(\text{CH}_2)_2$	Et	7-Me	EtOH	5f 43
<i>g</i>	$\text{CH}\equiv\text{C}(\text{CH}_2)_3$	Et	H	EtOH	5g 41
<i>h</i>	Et	Me	H	MeOH	5h 42
<i>i</i>	Et	Me	5-Br	MeOH	5i 41
<i>j</i>	$\text{MeOOC}(\text{CH}_2)_4$	Me	H	MeOH	5j 36
<i>k</i>	Et	Pr	H	PrOH	5k 46
<i>l</i>	Et	Pr	7-Me	PrOH	5l 45
<i>m</i>	$\text{CH}_3(\text{CH}_2)_6$	Pr	H	PrOH	5m 38

¹ Overall yield of the pure isolated product.

It is important to highlight that, in order to prevent the transesterification reaction at C-2, the indolization was conducted using the same alcohol (R^3OH) of the ester moiety. Detailed spectroscopic data for compounds 5a-m are included in the Supplementary Materials.

4. Conclusions

In conclusion, we disclosed a new, important application of β -nitroacrylates as a valuable alternative of α -ketoesters for synthesizing alkyl indole-2-carboxylates, which were key and useful intermediates of biologically active molecules. The method involved two easy steps and afforded title targets with a good substrate scope generality and satisfactory overall yields. Moreover, thanks to the possibility of removing the TMG by an easy filtration through the silica (first step), and the use of heterogeneous Amberlyst 15 in the second step, it was possible to avoid the typical tedious and wasteful aqueous work-up, with evident advantages from a sustainability point of view.

Supplementary Materials: The following are available online at <http://www.mdpi.com/2076-3417/9/23/5168/s1>, Spectroscopic Data.

Author Contributions: Conceptualization, S.G. and D.P.; methodology, S.G.; investigation, D.P.; validation, S.G. and D.P.; data curation, S.G.; writing—original draft preparation, A.P.; writing—review and editing, A.P.; supervision, R.B.; project administration, A.P.; funding acquisition, A.P.

Funding: This research received no external funding.

Acknowledgments: The authors thank the University of Camerino for financial support.

Conflicts of Interest: The authors declare no conflict of interest.

Abbreviations

The following abbreviations are used in this manuscript:

TMG	1,1,3,3-Tetramethylguanidine
DBU	1,5-Diazabicyclo[5.4.0]undec-5-ene
PS-TBD	1,5,7-Triazabicyclo[4.4.0]dec-5-ene bound to polystyrene (Sigma-Aldrich code: 01961)
PS-Carbonate	Carbonate on polymer support (Sigma-Aldrich code: 21850)
NNRTI	Non-nucleoside reverse transcriptase inhibitor
PPA	Poly Phosphoric Acid

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