

Cyclization Reactions of Homopropargyl Azide Derivatives Catalyzed by PtCl₄ in Ethanol Solution: Synthesis of Functionalized Pyrrole Derivatives

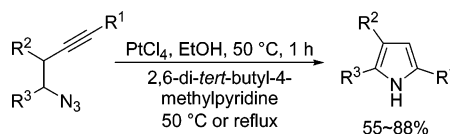
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ABSTRACT



PtCl₄-catalyzed cyclization reactions of homopropargyl azide derivatives to pyrrole rings were investigated. Using ethanol as solvent with 2,6-di-tert-butyl-4-methylpyridine as the base was found to be the best set of conditions for effecting this ring-closing reaction. These reaction conditions can be applied to the preparation of functionalized pyrrole derivatives, with no effect on the functional groups.

Carbon–nitrogen bond formation reactions are an important tool in the synthesis of biologically active heterocyclic substances.¹ In our efforts directed toward the development of new synthetic methods for obtaining heterocyclic compounds, we had previously reported effective methods for the synthesis of substituted indole derivatives, catalyzed by Cu(II) salts² or Pd(PPh₃)₄–methyl propiolate complex.³ Intermolecular nucleophilic addition reactions of the nitrogen atom to the alkyne moiety, that do not conjugate to the sp²- or sp-hybridized carbon, tend to be difficult, apparently due

to the low reactivity of the alkyne group. Until now, cyclization reactions, which involve the formation of carbon–nitrogen bonds, catalyzed by early transition metal or organolanthanoide complexes have been well investigated.^{1,4} In addition, examples for the same type reactions catalyzed by late transition metal complex, which contain, e.g., Pd, Au, Ag, Cu, and Ru, also have been reported.^{1,5} In spite of the powerful catalytic properties of Cu(II) salts² and Pd(PPh₃)₄–methyl propiolate complex,³ the cyclization reactions of **1** to **2** could not be effected, and starting material was recovered quantitatively (Scheme 1). After several attempts, we found that the cyclization reaction of **1** was

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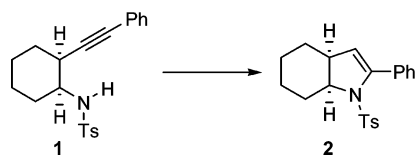
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Scheme 1. Cyclization Reactions of the Tosylamide 1 System



Reaction Conditions

A: Cu(OAc)₂ or Cu(OTf)₂

ClCH₂CH₂Cl, reflux: **No reaction**

B: Pd(PPh₃)₄, H—C≡C—CO₂Me, ZnBr₂, *i*-Pr₂NEt

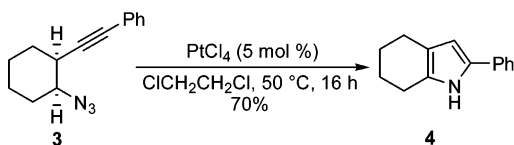
CH₂Cl₂ or THF, reflux: **No reaction**

C: PtCl₄ (5 mol %), CH₂Cl₂, reflux, 6.5 h:
68% yield

catalyzed by platinum catalyst, recently reported to be an activator of π -bonds (Scheme 1).^{6,7}

Disappointingly, it was found that the PtCl₄-catalyzed cyclization reaction can only be applied to a limited number of substrates, due to the instability of the enamine moiety of the products. However, in the process of surveying new substrates and catalyst systems, we unexpectedly found that homopropargyl azide derivatives could be converted to pyrrole rings in the presence of catalytic amounts of PtCl₄, in 70% yield (Scheme 2). While we were working on

Scheme 2. PtCl₄-Catalyzed Cyclization Reaction of Homopropargyl Azide Derivatives

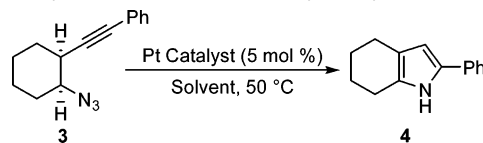


optimizing our reaction conditions, Toste's research group published a paper describing nearly identical reactions, including the conversion of homopropargyl azide derivatives to substituted pyrroles in the presence of catalytic amounts of (dppm)Au₂Cl₂ and AgSbF₆ in dichloromethane under mild conditions.⁸ Toste's group proposed a reaction mechanism in which gold(I) induces activation of the alkyne toward nucleophilic addition of the proximal nitrogen atom of the azide. Herein, we describe a simpler catalytic system for efficient pyrroles synthesis as Toste's reaction.⁹

Optimization of the Pt-catalyzed reactions began with the selection of the solvent. The long reaction time (16 h)

necessary in 1,2-dichloroethane was presumed to be due to low solubility of PtCl₄ (Table 1, entry 1). This problem was

Table 1. Cyclization Reactions Catalyzed by PtCl₄ or PtCl₂



entry	Pt catalyst	solvent	time (h)	yield (%)
1	PtCl ₄	ClCH ₂ CH ₂ Cl	16	70
2	PtCl ₄	EtOH	9	65
3 ^b	PtCl ₄	EtOH	1	79
4	PtCl ₄	<i>tert</i> -BuOH	24	2 (58) ^a
5	PtCl ₂	ClCH ₂ CH ₂ Cl	24	10 (76) ^a
6	PtCl ₂	EtOH	24	21 (67) ^a
7 ^b	PtCl ₂	EtOH	24	40 (48) ^a

^a The numbers in parentheses are yields of recovered **3**. ^b The reactions were started after stirring the platinum catalyst in ethanol for 1 h at 50 °C.

resolved by changing the solvent to ethanol (Table 1, entry 2). Surprisingly, the reaction rate was much improved by stirring the solution of PtCl₄ in ethanol for 1 h at 50 °C before adding the substrate (Table 1, entry 2 vs 3). The requirement of the induction period for obtaining higher catalytic activities suggested the possibility that initially added PtCl₄ is transformed to other active species.

In 1974, Hass and Hauthal reported that ethanol was oxidized by PtCl₄ to produce PtCl₂, acetaldehyde, and hydrogen chloride.¹⁰ Noting from our results that *tert*-butanol was not an effective solvent (Table 1, entry 4), we speculated that perhaps the active species in ethanol is PtCl₂, which could be generated in situ. However, when the reaction was performed in the presence of commercially available PtCl₂, completely different results were obtained than from the reaction with PtCl₄. Namely, the reaction rates were much slower in either dichloroethane or ethanol, and the rates could not be improved, even with an induction period (as in entry 3). Furthermore, the reaction did not proceed to completion even after 24 h (Table 1, entries 5, 6, and 7). From these results, it is obvious that PtCl₄ in ethanol is being converted into a species other than PtCl₂. However, the true active species cannot be identified at the present time. On the other hand, since it is difficult to imagine that Pt(IV) could be reduced to Pt(II) in dichloroethane, the structure of the catalyst may vary depending upon the solvent used.

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Table 2. Cyclization Reactions in the Presence of Base^a

entry	PtCl ₄ (mol %)	base (mol %)	yield (%)
1	5		50 (17) ^b
2	5	Cs ₂ CO ₃ (20)	0 (68) ^b
3	5	proton-sponge (20)	0 (91) ^b
4	5	2,6-di- <i>tert</i> -butyl-4-methylpyridine (20)	43 (44) ^b
5	15	2,6-di- <i>tert</i> -butyl-4-methylpyridine (60)	72

^a For all entries, **5** and the base were added after the solution of PtCl₄ in ethanol was stirring for 1 h at 50 °C. ^b The numbers in parentheses are yields of recovered **5**.

Since pyrroles are π -electron-rich heteroaromatic compounds, it is known that the pyrrole ring may be prone to

decomposition even under weakly acidic conditions. In fact, when the reaction was applied to **5**, the pyrrole **6** gradually decomposed over the long course of the reaction time. Because the possibility for generation of hydrogen chloride could not be ruled out, we next ran the reaction in the presence of a variety of bases. The addition of either the inorganic base Cs₂CO₃ or the proton-sponge completely inhibits the reaction (Table 2, entries 2 and 3). On the other hand, in the presence of a bulkier base, such as 2,6-di-*tert*-butyl-4-methylpyridine, the stoichiometry of the reaction was much improved over that obtained in the reaction run in the absence of base [Table 2, entry 1 (**5** + **6** = 67%) vs 4 (**5** + **6** = 87%)], although the reaction rate was slightly suppressed. Finally, the reaction could be run to completion by increasing the amount of both PtCl₄ and the base to provide the pyrrole **6** in 72% yield.

Having established the reaction conditions, we next applied it to other substrates with a variety of functional groups, producing the results compiled in Table 3. The reaction conditions have to be modified slightly, depending on the

Table 3. Platinum-Catalyzed Cyclization Reactions^g

entry	homopropargyl azide	product	T °C	yield (%) (time) under Ar atmospher	under aerobic condition
1 ^a			50 °C	88 (4 h)	78 (60 min)
2 ^b			50 °C	72 (120 min)	—
3 ^b			50 °C	55 (90 min)	24 (80 min)
4 ^b			50 °C	65 (60 min)	—
5 ^b			50 °C	64 (60 min)	—
6 ^c			50 °C	69 (30 min)	23 (14) ^f (60 min)
7 ^d			reflux	74 (6 h)	70 (3) ^f (160 min)
8 ^d			reflux	61 (14 h)	—
9 ^d			reflux	57 (82) ^e (50 min)	65 (31) ^f (80 min)

^a PtCl₄ (5 mol %) and 2,6-di-*tert*-butyl-4-methylpyridine (20 mol %) in 0.1 M substrate concentration in ethanol. ^b PtCl₄ (15 mol %) and 2,6-di-*tert*-butyl-4-methylpyridine (60 mol %) in 0.1 M substrate concentration in ethanol. ^c PtCl₄ (5 mol %) and 2,6-di-*tert*-butyl-4-methylpyridine (20 mol %) in 0.03 M substrate concentration in ethanol. ^d PtCl₄ (15 mol %) and 2,6-di-*tert*-butyl-4-methylpyridine (60 mol %) in 0.01 M substrate concentration in ethanol. ^e The number in parentheses was the yield of the reaction in 1,2-dichloroethane. ^f The number in parentheses was the yield of the recovered starting material. ^g For all entries, both substrate and base were added after stirring the solution of PtCl₄ in ethanol for 1 h at 50 °C.

structure of the substrate, in order to obtain maximum chemical yield. Namely, the substrate bearing the phenylethynyl group is the most reactive and just 5 mol % of PtCl₄ and 20 mol % of 2,6-di-*tert*-butyl-4-methylpyridine are enough to afford the product in good yield (Table 3, entry 1). Substrates with an alkyl group substituted on the alkyne terminal (Table 3, entry 3), as well as compounds with silyl ether (Table 3, entry 4), hydroxyl (Table 3, entry 2), and ethoxycarbonyl groups (Table 3, entry 5), gave the corresponding pyrrole rings in reasonable yields under standard reaction conditions (15 mol % of PtCl₄ and 60 mol % of 2,6-di-*tert*-butyl-4-methylpyridine, 0.1 M solution of the azide in ethanol). The amino group in the substrate must be protected in order to obstruct complexation between the amino group and PtCl₄. In our case, *N*-Boc tosylamide was used as the protecting group; however, partial elimination of the Boc group in the product was observed under standard reaction conditions. This side reaction was avoided by diluting the reaction mixture from 0.1 to 0.03 M solution (Table 3, entry 6).

It is known that terminal alkynes react with organic azides to produce triazole compounds.¹¹ In our reactions, we observed low yields of the products under standard reaction conditions, presumably due to intermolecular polymerization

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(12) The reaction of (4-azidobutyl)benzene in the presence of 15 mol % of PtCl₄ and 60 mol % of 2,6-di-*tert*-butyl-4-methylpyridine, which was stirred in EtOH for 1 h at 50 °C before adding the substrate, at 50 °C gave the starting material in 80% yield and the complete decomposition of the starting material was observed in the presence of 100 mol % of PtCl₄ and 2,6-di-*tert*-butyl-4-methylpyridine.

of the acyclic starting materials. However, diluting the reaction conditions proved effective for such substrates. In these cases, 0.01 M proved to be the best concentration for the substrate, and the corresponding pyrroles **16**, **18**, and **20** were afforded in reasonable yields (Table 3, entries 7, 8, and 9).

Importantly, these reaction conditions can be carried out under aerobic condition for phenyl-substituted pyrrole syntheses (Table 3, entries 1, 7, and 9). For these substrates, the solvent (ethanol) also can be used without any purification. However, decomposition of the pyrroles during the reactions for the other substrates was observed (Table 3, entries 3 and 6).

One possibility for the mechanism of this cyclization reaction may be the same as that proposed by Toste.⁸ However, we observed that the azide group was completely decomposed under the cyclization reaction conditions.¹² Thus, another pathway cannot be ruled out. Clarification of the mechanism, further improvement of the conditions, and applications to the synthesis of biologically active pyrrole-containing compounds are currently underway in our laboratory.

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Supporting Information Available: Experimental detail for the synthesis of the homopropargyl azide derivatives, a general experimental procedure for the cyclization reaction, and spectroscopic data for all new compounds. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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