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Stereoselective preparation of conjugated *E*-enynes from *E*-vinylic tellurides and terminal alkynes *via* Sonogashira cross-coupling[†]

Gilson Zeni,*^{*a*} Diego Alves,^{*a*} Jesus M. Pena,^{*a*} Antonio L. Braga,^{*a*} Helio A. Stefani^{*b*} and Cristina W. Nogueira^{*a*}

^a Departamento de Química, Laboratório de Bioquímica Toxicológica, UFSM, Santa Maria, RS, Brazil 97105-900. E-mail: gzeni@quimica.ufsm.br

^b Faculdade de Ciências Farmacêuticas, USP, São Paulo, SP, Brazil

Received 23rd January 2004, Accepted 4th February 2004 First published as an Advance Article on the web 9th February 2004

E-Vinylic tellurides undergo a direct cross-coupling reaction with terminal alkynes in the presence of palladium(II)/ CuI in Et₃N at room temperature to give *E*-enynes in good yields. The methodology represents a general and efficient protocol for carrying out the synthesis of *E*-enynes under mild conditions with complete retention of configuration.

Vinylic tellurides are useful intermediates in organic synthesis.¹ Of the two isomers, the Z-vinylic tellurides have been employed more frequently as intermediates because of easy availability of these species.² One of the most important reactions of vinylic tellurides is their transmetallation.³ The vinyl organometallic obtained in this way can react with carbonyl compounds,⁴ α , β -unsaturated systems⁵ or epoxides.⁶ Lately, we have employed vinylic tellurides in the synthesis of polyacetylenic acids isolated from *Heisteria acuminata* by using a vinylic telluride cross-coupling reaction, Scheme 1.⁷



Recently, new applications of vinylic tellurides utilizing palladium catalyzed cross-coupling have been described.⁸ In this case, they act as an equivalent of an aryl or vinyl carbocation. They react in a similar way to vinylic halides or triflates in the Sonogashira,⁹ Heck,¹⁰ Suzuki¹¹ and Stille¹² reactions. Our continuous efforts in the synthesis of enynes by cross-coupling reaction¹³ led us to find out that the *E*-vinylic tellurides **1a–c** can be effectively applied to the synthesis of *E*-enynes *via* a cross-coupling reaction with terminal alkynes, using palladium as catalyst, Scheme 2.¹⁴



Scheme 2

The starting materials required for the coupling, *E*-vinylic tellurides **1a–c**, were synthesized by stereospecific *cis* hydroalumination of the acetylene,¹⁵ followed by transmetallation of the *E*-vinyl aluminate complexes formed with organotellurenyl bromide/LiCl¹⁶ (Scheme 3).

† Electronic supplementary information (ESI) available: spectroscopic data for all new compounds as well as detailed experimental procedures. See http://www.rsc.org/suppdata/ob/b4/b401059k/

 Table 1
 Influence of the catalyst in the cross-coupling reaction

Entry	Catalyst (mol%)	Time (h)	Yield, 2a ^{<i>a</i>} (%)
1	Pd(PPh ₃)₄/CuI (20)	48	0
2	$Pd(PPh_3)_4$ (20)	48	0
3	$PdCl_2/PPh_3$ (20)	20	8
4	$PdCl_{2}(PPh_{3})_{2}$ (20)	20	5
5	Pd(OAc), (20)	24	2
6	$PdCl_2(PhCN)_2(20)$	20	5
7	PdCl ₂ /CuI (1)	24	23
8	PdCl ₂ /CuI (3)	24	30
9	PdCl ₂ /CuI (5)	24	47
10	PdCl ₂ /CuI (10)	12	62
11	PdCl ₂ /CuI (20)	3	76
12	$PdCl_2(20)$	24	28

^{*a*} All reactions were carried out with 1 mmol of *E*-vinylic tellurides **1a**, 2 mmol of 2-propyn-1-ol in methanol (5 mL) at room temperature in the presence of palladium catalyst and Et_3N (0.8 mL). Entries 1 and 7–11 used CuI.



Our initial efforts were focused on the reactivity of *E*-vinylic telluride 1c in the cross coupling reaction with 1-alkynes. Thus, 1c (1 equiv.) was treated in methanol at room temperature with 2-propyn-1-ol (2 equiv.) in the presence of $PdCl_2$ (20 mol%)/CuI (20 mol%), and Et_3N (0.8 mL) as the base. Under these conditions, the corresponding enyne 2a was obtained as a minor product, (Scheme 4). This may be explained by the possible transmetallation of Te with Pd at both the vinylic and aryl carbon giving undesirable homocoupling products.



We tried reacting *E*-vinylic telluride 1c in methanol at room temperature with 2-propyn-1-ol in the presence of PdCl₂ as catalyst, and Et₃N as base, in the absence of CuI. Under these conditions, the cross-coupling reaction still proceeded unsatisfactorily.

In view of these disappointing results, we decided to explore the reaction with the *E*-vinylic tellurides **1a** and **1b** under the same experimental conditions described above. In the course of the study, we found that the cross-coupling reaction of *E*-vinylic

 Table 2
 E-Enynes 2 Prepared According to Scheme 2¹⁷

Entr	y Telluride 1	<i>E</i> -Enyne 2	Time (h)	Yield (%)
1	PhTe(<i>n</i> -Bu)	Ph2a	3	76
2	1a	Ph2b	5	67
3	1a	Ph2c	6	68
4	1a	HO Ph 2d	5	70
5	1a	Ph2e	6	75
6	<i>n</i> -Bu 1b Te(<i>n</i> -Bu)	Ph n-Bu2f	3	78
7	1b	n-Bu	6	75
8	1b	HO n-Bu 2h	5	64
9	1b	n-Bu 2i	4	73
10	1b	n-Bu	3	70
		Ph		

tellurides 1a and 1b with 1-alkynes are best catalyzed by PdCl₂ (20 mol%)/CuI (20 mol% each) (entry 11; Table 1). Thus, the reaction of 1a (1 equiv.) with 2-propyn-1-ol (2 equiv.) over 3 h in methanol at room temperature using PdCl₂/CuI (20 mol%) gave the E-enyne, in 76% yield. Other palladium complexes such as Pd(PPh₃)₄/CuI, Pd(PPh₃)₄, PdCl₂/PPh₃, PdCl₂(PPh₃)₂, Pd(OAc)₂, PdCl₂(PhCN)₂ were less effective (Table 1). The reaction was greatly enhanced by increasing the amount of catalyst from 1% to 20% (entries 7-11). A rather different trend was noted when the reaction was performed in the absence of copper salts, such as CuI, (entry 12; Table 1). This reaction is also highly affected by the solvent. The performance (yield of enyne) of various solvents in the same reaction under identical conditions decreased as follow [PdCl2 (20 mol%)/CuI (20 mol%), rt, 3 h]: THF, acetonitrile, DMF (40-55%); pyridine, DME (22-35%); toluene, CH₂Cl₂, 1,2-dichloroethane, hexane (no cross-coupling product was observed).

The nature of the amine was critical for the success of the coupling. When different amines (piperidine, pyrrolidine, n-BuNH₂, Et₂NH, i-Pr₂NH) were used in lieu of Et₃N, unsatisfactory yields of the desired enynes were obtained. Careful analysis of the optimized reactions revealed that the optimum conditions for the coupling in Scheme 2 were found to be the

use of PdCl₂ (20 mol%)/CuI (20 mol%), MeOH (5 mL), (*E*)vinylic telluride **1a** or **1b** (1 mmol), the appropriate alkyne (2 mmol) and Et₃N (0.8 mL) at 25 °C. Extending the coupling reaction to other alkynes, various *E*-enynes **2a**–**j** were obtained in good yields (Table 2).¹⁷

The formation of enynes was confirmed by the analysis of the ¹H NMR spectra. The stereoisomeric purities of the *E*-enynes **2a**–**j** were similar to starting *E*-vinylic tellurides **1a**–**b**,¹⁴ due to a complete retention of configuration in this type of reaction. The stereochemistry of the obtained enynes was easily established. As an example, the ¹H NMR spectrum of compound **2a**, showed a doublet at 6.8 ppm with a coupling constant of 16.3 Hz and a double triplet at 6.0 ppm with coupling constants of 16.3 Hz and 1.9 Hz. These coupling constants confirm the *trans* relationship of the hydrogen of the double bond.

In summary, we have explored the Pd(II) catalyzed crosscoupling reaction of *E*-vinylic tellurides with alkynes and established a new stereoselective route to *E*-enynes in good yields. The reaction proceeds cleanly under mild conditions. The cross coupling reaction was sensitive to the nature of the amine, catalyst and solvent. This novel approach to functionalized enynes could open a new route to the biologically important

Acknowledgements

We are grateful to the FAPERGS, CAPES, CNPq, UFSM and FAPESP (98/10821–0) for financial support. CNPq is also acknowledged for Ms (D. A.) fellowship.

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- 16 M. J. Dabdoub and T. M. Cassol, *Tetrahedron*, 1995, **51**, 12971– 12982.
- 17 Typical procedure for the preparation of enynes 2: To a two-necked 25 mL round-bottomed flask under an argon atmosphere containing PdCl2 (0.036 g, 20 mol%), CuI (0.038 g, 20 mol%) and dry methanol (5 mL) was added Z-vinylic tellurides 1a (0.287 g, 1 mmol). After stirring the mixture for 15 minutes at room temperature, 2-propyn-1ol (0.112 g, 2 mmol) and Et₃N (0.8 mL) were added. The reaction was stirred at room temperature for 3 h. After this time the solid part was filtered under vacuum and the filtrate was treated with saturated brine and extracted with dichloromethane $(3 \times 25 \text{ mL})$. The combined organic layers were dried over MgSO4 and concentrated under vacuum. The residue was purified by flash chromatography and eluted with hexane/ethyl acetate (80 : 20). Selected spectral and analytical data for 2a: (4E)-5-phenylpent-4-en-2-yn-1-ol: yield 0.120 g (76%); ¹H NMR: CDCl₃, 200 MHz, δ(ppm): 7.21-7.15 (m, 5H), 6.81 (d, J = 16.3 Hz; 1H), 6.00 (dt, J = 16.3 and 1.9 Hz; 1H), 4.32 (s; 2H), 2.45 (s, 1H); ¹³C NMR: CDCl₃, 50 MHz, δ (ppm): 141.76, 135.88, 128.60, 126.20, 107.38, 89.39, 84.74, 51.46. MS m/z (%) 158 (55), 140 (50), 129 (100), 115 (52), 102 (12), 77 (15), 51 (12).