



Decarboxylative and dehydrative coupling of dienoic acids and pentadienyl alcohols to form 1,3,6,8-tetraenes

Ghina'a I. Abu Deiab^{‡1}, Mohammed H. Al-Huniti^{‡1}, I. F. Dempsey Hyatt^{1,§}, Emma E. Nagy¹, Kristen E. Gettys¹, Sommayah S. Sayed¹, Christine M. Joliat¹, Paige E. Daniel¹, Rupa M. Vummalaneni¹, Andrew T. Morehead Jr², Andrew L. Sargent² and Mitchell P. Croatt^{*1}

Full Research Paper

[Open Access](#)**Address:**

¹Department of Chemistry and Biochemistry, University of North Carolina at Greensboro, Greensboro, NC 27402, USA and
²Department of Chemistry, East Carolina University, Greenville, NC 27858, USA

Email:

Mitchell P. Croatt* - mpcroatt@uncg.edu

* Corresponding author ‡ Equal contributors

§ Current address: Department of Chemistry, Adelphi University, Garden City, NY 11530, USA

Keywords:

decarboxylation; diene; dienolate; palladium; pentadienyl; tetraene

Beilstein J. Org. Chem. **2017**, *13*, 384–392.

doi:10.3762/bjoc.13.41

Received: 17 November 2016

Accepted: 03 February 2017

Published: 28 February 2017

Associate Editor: B. Stoltz

© 2017 Deiab et al.; licensee Beilstein-Institut.

License and terms: see end of document.

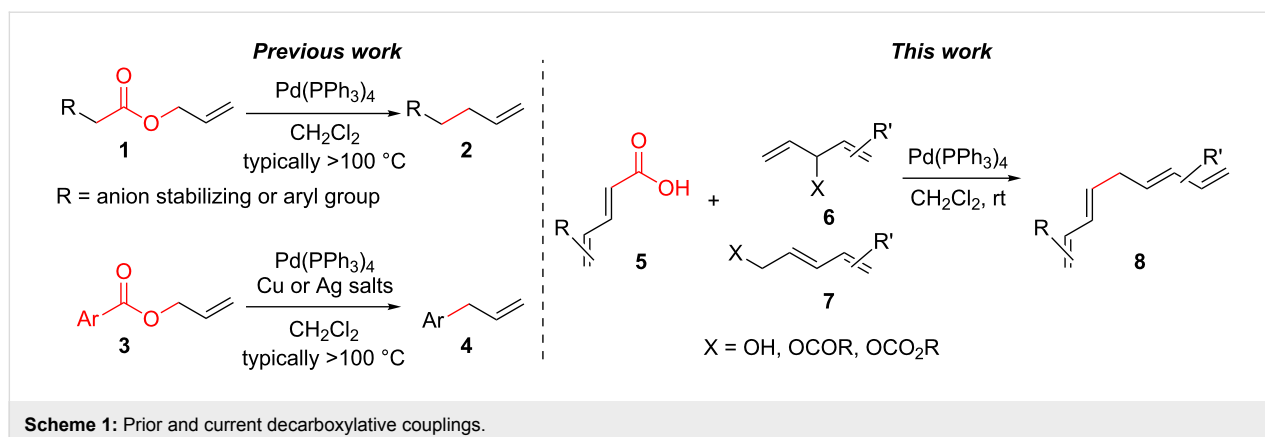
Abstract

Dienoic acids and pentadienyl alcohols are coupled in a decarboxylative and dehydrative manner at ambient temperature using Pd(0) catalysis to generate 1,3,6,8-tetraenes. Contrary to related decarboxylative coupling reactions, an anion-stabilizing group is not required adjacent to the carboxyl group. Of mechanistic importance, it appears that both the diene of the acid and the diene of the alcohol are required for this reaction. To further understand this reaction, substitutions at every unique position of both coupling partners was examined and two potential mechanisms are presented.

Introduction

The construction of sp^2 – sp^3 carbon–carbon bonds remains a difficult and important problem in organic synthesis. Cross-coupling reactions provide avenues to these otherwise difficult reactions, but often require prefunctionalization of the coupling partners [1–9]. However, recent C–H activation research has enabled the use of further simplified starting materials [10–18].

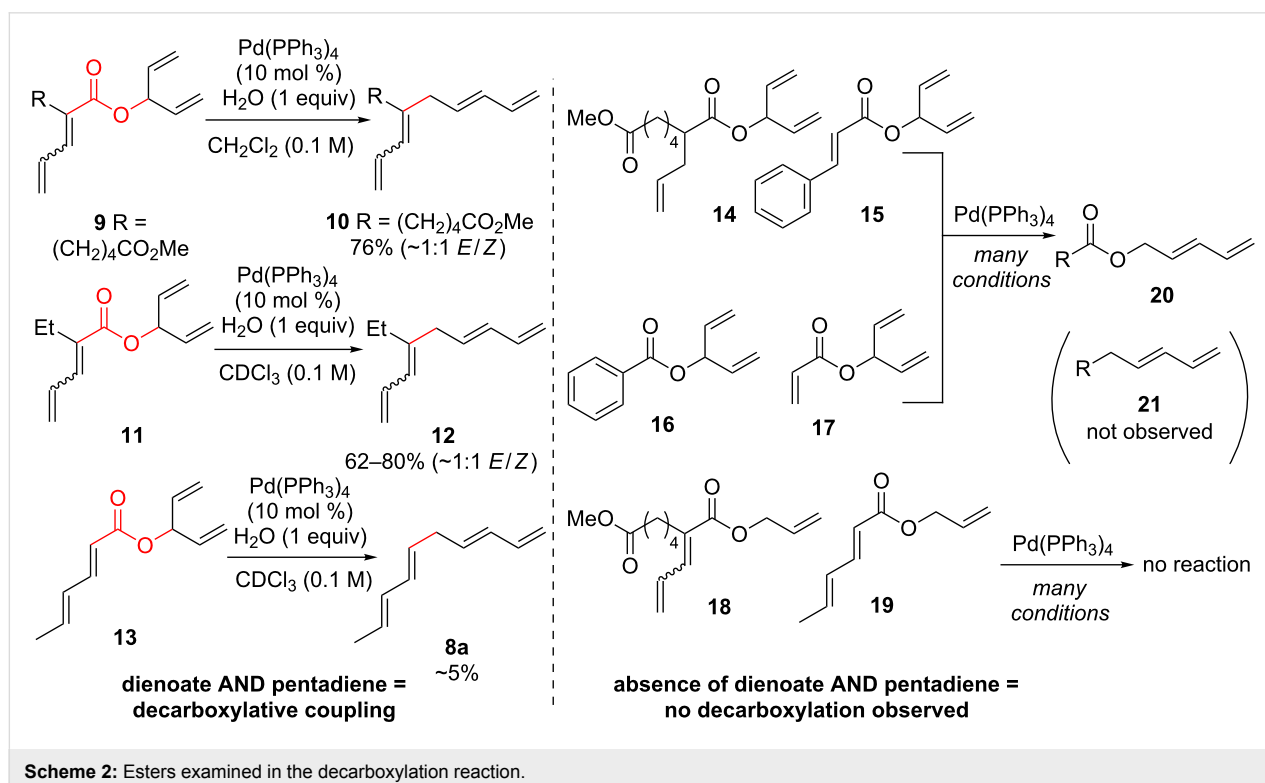
Another approach to the formation of C–C bonds is through decarboxylative coupling reactions (Scheme 1). This can be arrived in a one-component fashion via the removal of CO_2 from an ester or in a two-component manner by removal of CO_2 from a carboxylic acid and coupling this to a substrate with a benzylic or allylic leaving group [19,20].



Typical Pd(0)-catalyzed decarboxylative coupling reactions utilize an allylic or benzylic ester with either an anion-stabilizing group adjacent to the carboxyl group (i.e., carbonyl [19,21,22], nitrile [23–25], nitro [26,27], or alkyne [21,28–32], Scheme 1), or use an aryl carboxylate [33,34] which typically requires the assistance of silver or copper(I) salts for the decarboxylative step. It is rare to use a pentadienyl electrophile [35], or to have a diene or simple alkene adjacent to the carboxyl group [20,36–39]. Despite the absence of this type of reactivity, the decarboxylative coupling of a pentadienyl dienoate (**9**; Scheme 2) was desirable enough for our group's synthesis of clinprost that we attempted the reaction [40,41]. Fortunately, this coupling reaction was successfully employed

in our reported nine-step synthesis of clinprost [41]. A structurally related compound (**11**) reacted similarly, however, the sorbate derivative (**13**) was low yielding with the majority of the material only rearranging to the linear ester. In all three of these cases, we never observed the more stable, fully conjugated tetraene. "Skipped diene" motifs are found in various natural products and there are few methods available to prepare these dienes [42–52]. Skipped tetraene systems have even fewer methods for their synthesis [53–55], which makes the method described herein even more valuable.

It was determined that modifying the dienoate motif yielded only the rearranged product under the reaction conditions, in-



cluding the dihydro (**14**), cinnamate (**15**), benzoate (**16**), and acrylate (**17**) analogues (Scheme 2). Moreover, allylic dienoates **18** and **19** gave no reaction with Pd(0) catalysis. These results led us to the determination that there was a unique reactivity imbued to the molecule by having both the dienoate and penta-dienyl moieties. Herein, are presented more details for this reaction, including the substrate scope for the intermolecular case.

Results and Discussion

In addition to determining the requisite nature of both the penta-dienyl and dienoate groups, it was found that trace amounts of water were required for decarboxylative coupling (Table 1). For example, careful exclusion of water from reagents and solvent and performing the reaction in the glovebox led to formation of rearranged product and no decarboxylative coupling reaction (**22**, Table 1, entry 1). Less than 1 equivalent of water allowed

for a slow reaction and incomplete conversion, 1–2 equivalents was optimal with yields around 70% and more water was not beneficial (Table 1, entries 2–8). The use of equimolar amounts of methanol and water as a proton source allowed for decarboxylation to take place but with a low yield (Table 1, entry 9) and the reaction run in TFE as a solvent did not result in any decarboxylation (Table 1, entry 10).

In addition to the requirement for water, it was determined that phosphine ligands were necessary (Table 1, entry 11), either as ligands or as participants in the reaction as discussed later. The typical catalyst used, Pd(PPh₃)₄, worked well, however, it was found that a more ideal ratio of palladium metal to ligand was 1:1 or 1:2, with greater amounts of triphenylphosphine lowering the reaction yield when using the Pd₂dba₃ catalyst (Table 1, entries 12–14). It was determined that reactions performed in

Table 1: Optimization of the one-component decarboxylation reaction.^a

Entry	Catalyst	Solvent	Additives	Yield of 10 (Yield of 22) ^b
1	Pd(PPh ₃) ₄	CH ₂ Cl ₂	anhydrous	0% (99%)
2	Pd(PPh ₃) ₄	CH ₂ Cl ₂	0.5 equiv H ₂ O	27%
3	Pd(PPh ₃) ₄	CH ₂ Cl ₂	1.1 equiv H ₂ O	77%
4	Pd(PPh ₃) ₄	CH ₂ Cl ₂	1.3 equiv H ₂ O	72%
5	Pd(PPh ₃) ₄	CH ₂ Cl ₂	silylated glass, 1 equiv H ₂ O	55% (15%)
6	Pd(PPh ₃) ₄	CH ₂ Cl ₂	dry glass balls	37% (24%)
7	Pd(PPh ₃) ₄	CH ₂ Cl ₂	wet glass balls	51%
8	Pd(PPh ₃) ₄	CH ₂ Cl ₂ /H ₂ O	biphasic	49%
9	Pd(PPh ₃) ₄	CH ₂ Cl ₂	1 equiv MeOH, 1 equiv H ₂ O	33% (26%)
10	Pd(PPh ₃) ₄	TFE	trace CH ₂ Cl ₂	0%
11	Pd ₂ (dba) ₃	CH ₂ Cl ₂	0 mol % PPh ₃ , 1 equiv H ₂ O	0%
12	Pd ₂ (dba) ₃	CH ₂ Cl ₂	10 mol % PPh ₃ , 1 equiv H ₂ O	64%
13	Pd ₂ (dba) ₃	CH ₂ Cl ₂	20 mol % PPh ₃ , 1 equiv H ₂ O	61%
14	Pd ₂ (dba) ₃	CH ₂ Cl ₂	30 mol % PPh ₃ , 1 equiv H ₂ O	12%
15	Pd ₂ (dba) ₃	CH ₂ Cl ₂	10 mol % L1 , 1 equiv H ₂ O	70%
16	Pd ₂ (dba) ₃	CH ₂ Cl ₂	10 mol % L2 , 1 equiv H ₂ O	70%
17	Pd ₂ (dba) ₃	CH ₂ Cl ₂	10 mol % L4 , 1 equiv H ₂ O	18%
18	Pd ₂ (dba) ₃	CH ₂ Cl ₂	10 mol % L5 , 1 equiv H ₂ O	10%
19	Pd(OAc) ₂	CH ₂ Cl ₂	1 equiv H ₂ O	0%
20	Pd(OAc) ₂	CH ₂ Cl ₂	40 mol % PPh ₃ , 1 equiv H ₂ O	10%
21	none	CH ₂ Cl ₂	1 equiv PPh ₃ , 1 equiv H ₂ O	0%

^aReaction Conditions: Pd metal (10 mol %) and the indicated solvent and additives for 24 hours. ^bIsolated yields.

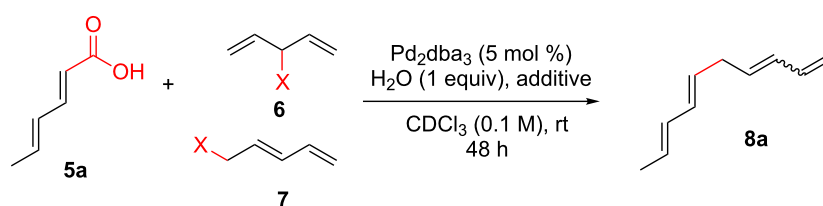
the presence of electron-rich ligands had both quicker kinetics and more efficient yields compared to electron-deficient ligands (Table 1, entries 15–18 and Supporting Information File 1 for kinetic information). Although not as efficient, it was found that a palladium(II) catalyst functioned in this reaction, presumably functioning as a pre-catalyst and being reduced in situ to the palladium(0) catalyst (Table 1, entries 19 and 20). As a control reaction, it was found that no reaction occurred in the absence of palladium catalyst (Table 1, entry 21).

As shown earlier, bis-allylic sorbate **13** (Scheme 2) was found to be low yielding for the decarboxylative coupling reaction. Reactions of sorbate **13** monitored by ^1H NMR showed nearly quantitative isomerization of the bis-allylic group into a linear pentadienyl system. Increasing the reaction time did not result in greater conversion to tetraene **8a**, which indicates that the products may be competitively ligating and poisoning the Pd(0) catalyst (see Supporting Information File 1 for additional evidence of product inhibition). The isomerization reaction to form **22** was presumably occurring via ionization of the allylic system using Pd(0), followed by recombination of the carboxylate at the terminal position of the pentadienyl system. Based on these data, we hypothesized that a two-component reaction using a dienolic acid and bis-allylic acetate might be possible, however, the presence of both water and a carboxylic acid would increase the possibility for isomerization of the 1,3,6,8-tetraenes into the fully conjugated 1,3,5,7-tetraenes, or possibly polymerization.

Despite the low yield for decarboxylation with sorbate **13**, the initial attempt used inexpensive sorbic acid as the dienolic acid. Gratifyingly, this reaction was successful and it was again determined that no isomerization to the fully conjugated system was observed (Table 2, entry 1). Other bis-allylic leaving groups were studied and, unexpectedly, it was determined that divinylcarbinol was superior (Table 2, entries 1–6). In fact, the better leaving groups were either slow or ineffective. This could be due to the less basic leaving groups not sufficiently deprotonating sorbic acid, which may be required for this reaction as is discussed mechanistically later (Scheme 3). Similar to the single component reaction, more than two equivalents of phosphine, relative to palladium metal, was detrimental (compare entries 12–14 of Table 1 with entries 6–8 of Table 2), however, the reaction was successful using Pd(PPh₃)₄ (Table 1, entry 9).

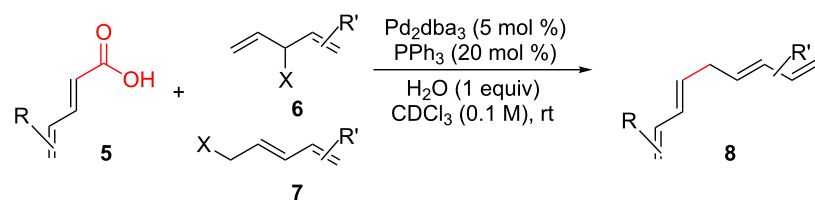
To further understand this interesting decarboxylative coupling reaction, a handful of different pentadienyl electrophiles and dienolic acids were examined (Table 3). Typically, the pentadienyl alcohol was used; however, in some cases the acetate was superior. It was found that both a methyl or phenyl substituent on the alcohol derivative would result in branched product **8b** or **8d** as a major product with a product ratio of 6:1 or 4:1, respectively (Table 3, entries 2–4). The yields for these reactions were low, but the remaining material was typically starting material and the ester where the acid and alcohol are coupled together. There was no effect on the yields upon leaving the reactions longer than 48 hours and it was found that the addition of

Table 2: Optimization of the two-component decarboxylation reaction.^a



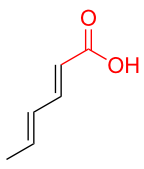
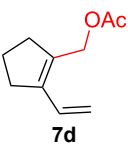
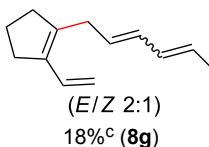
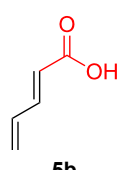
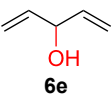
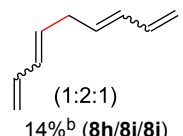
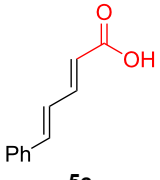
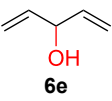
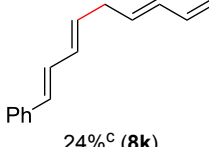
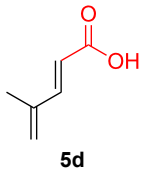
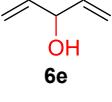
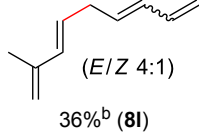
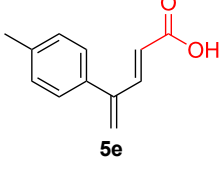
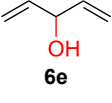
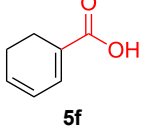
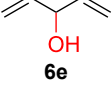
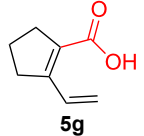
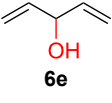
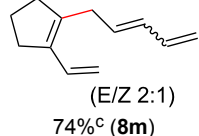
Entry	Pentadienyl group	Additive	Yield ^b
1	6a , X = OAc	PPh ₃ (20 mol %)	12%
2	6b , X = OCO ₂ Me	PPh ₃ (20 mol %)	35%
3	6c , X = OBz	PPh ₃ (20 mol %)	11%
4	6d , X = O ₂ C(4-CF ₃ Ph)	PPh ₃ (20 mol %)	6%
5	7a , X = Br	PPh ₃ (20 mol %)	0%
6	6e , X = OH	PPh ₃ (20 mol %)	40%
7	6e , X = OH	PPh ₃ (10 mol %)	18%
8	6e , X = OH	PPh ₃ (30 mol %)	24%
9 ^c	6e , X = OH	NA	28%

^aReaction conditions: Sorbic acid (**5a**, 1 equiv), pentadienyl group (**6** or **7**, 1 equiv), Pd₂(dba)₃·CHCl₃ (5 mol %) unless indicated otherwise, H₂O (1 equiv), in CDCl₃ for 48 hours. ^bNMR yields. ^cPd(PPh₃)₄ (10 mol %).

Table 3: Substrate scope for the two-component decarboxylation reaction.^a

Entry	Dienoic acid	Pentadienyl group	Yield (product) ^b
1			 (<i>E/Z</i> 10:1) 40% ^b (8a)
2			 (6:1) 8% ^c (8b/8c)
3			 (2:1) 21% ^b , 17% ^c (8d/8e)
4			 (4:1) 13% ^b , 6% ^c (8d/8e)
5			 16% ^c (8f)

Table 3: Substrate scope for the two-component decarboxylation reaction.^a (continued)

6	 5a	 7d	 (<i>E/Z</i> 2:1) 18% ^c (8g)
7	 5b	 6e	 (1:2:1) 14% ^b (8h/8i/8j)
8	 5c	 6e	 24% ^c (8k)
9	 5d	 6e	 (<i>E/Z</i> 4:1) 36% ^b (8l)
10	 5e	 6e	decomposition
11	 5f	 6e	decomposition
12	 5g	 6e	 (<i>E/Z</i> 2:1) 74% ^c (8m)

^aReaction conditions: Dienoic acid (**5**, 1 equiv), pentadienyl group (**6** or **7**, 1 equiv), H₂O (1 equiv), Pd₂(dba)₃·CHCl₃ (5 mol %), PPh₃ (20 mol %), in CDCl₃ for 48 h. ^bNMR yields due to volatility of product. ^cIsolated yields.

tetraene product inhibited the reaction (see Supporting Information File 1 for details). With these highly unsubstituted tetraene products, it is hypothesized that the product may be sequestering the palladium catalyst. Two cyclic dieny acetates were also studied (Table 3, entries 5 and 6) and they yielded tetraenes **8f** and **8g**. The dienes of entries 5 and 6 could have formed additional isomers by coupling to the other end of the pentadienyl group, but only one regioisomer was observed.

With respect to the dienoic acid, it was determined that the unsubstituted compound, pentadienoic acid, underwent decarboxylative coupling, although as a mixture of *E/E*, *E/Z*, and *Z/Z* isomers (Table 3, entry 7). Alkyl and aryl substituents were possible on the dienoate with the exception of an aryl group at the gamma position (Table 3, entries 8–10). Two cyclic dienoic acids were synthesized [56,57] and while the cyclohexadienoic acid did not decarboxylate (Table 3, entry 11), the vinylic-

clopentoic acid had a good yield of a complex tetraene (Table 3, entry 12).

Based on the information obtained during optimization and screening of compounds, two potential mechanisms are proposed (Scheme 3). Both options allow for the one (**13**) or two (**5** and **6**) component process to be used while also allowing for the reversible formation of linear ester **23**. Pathway B involves a Morita–Baylis–Hillman type process. The role of water would be to hydrogen bond to the carboxylate to make the system more electrophilic (**B**). This would accelerate the addition of the phosphine to generate zwitterion **C** [58]. Preliminary modeling for this ion indicates that both the electrophilic terminal vinyl group of the pentadienyl ligand and the nucleophilic α -carbon are in close proximity to one another. Formation of the carbon–carbon bond would then regenerate the Pd(0) catalyst and phosphonium carboxylate **D**. Decarboxylative elimination of the phosphine results in formation the 1,3,6,8-tetraene. It is proposed for pathway B that the dienophile is required so that the α -carbon is not blocked by the bulky phosphine group since it can add in a 1,6- or 1,4-manner, both reversibly.

Alternatively, pathway A has the palladium catalyst coordinate to one of the alkenes of the dienoate instead of the carboxylate (**E**). It is proposed that a water cluster around the carboxylate would enable this process by hydrogen bonding to the carboxylate. The conversion of **E** to **F** would form the C–C bond by having the palladium catalyst convert from one type of η^3 -allyl

and π -complex (**E**) to a different allyl/ π -complex (**F**). Finally, decarboxylative reduction of the palladium would release the product while regenerating the catalyst. Preliminary computational calculations using NEB [59] support pathway A and the HOMO of the transition state between **E** and **F** (Figure 1) calculated using the Gaussian 09 implementation of DFT with a B3LYP functional, 6-31g* basis, and polarized continuum model of solvation for DCM, shows close proximity of two in-phase orbitals for the requisite C–C bond, whereas removal

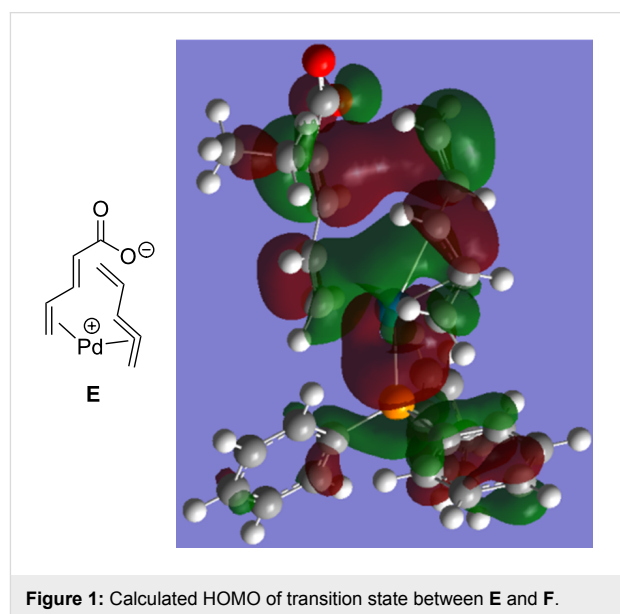
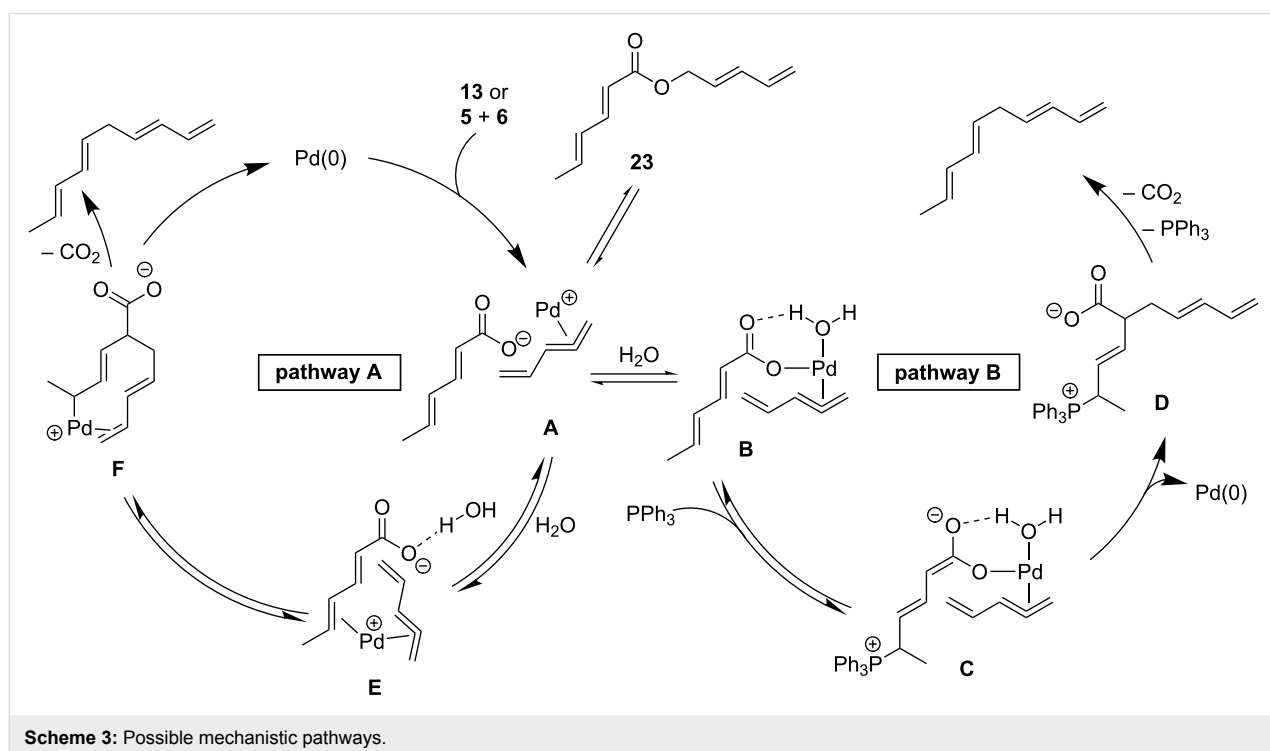


Figure 1: Calculated HOMO of transition state between **E** and **F**.



Scheme 3: Possible mechanistic pathways.

of any one of the alkenes from this structure would lead to anti-bonding relationships to bond to the alpha carbon.

Conclusion

In summary, we present information that is of value to advancing the area of metal-catalyzed decarboxylative coupling reactions, specifically those of pentadienyl dienoates that do not require an anion-stabilizing group, are run at ambient temperature, and can utilize the more accessible alcohol for a leaving group. This reaction was advanced to be possible in a two-component fashion, allowing for the conversion of dienoic acids and pentadienyl alcohols into 1,3,6,8-tetraenes with the only stoichiometric byproducts being water and carbon dioxide. These reactions currently require a diene motif with each coupling partner, but the product maintains the independent reactivity opportunities of these isolated dienes as opposed to forming the fully conjugated 1,3,5,7-tetraene. A variety of substrates were explored where each of the unique positions on the coupling partners was modified and two different mechanistic pathways are presented. A more in-depth mechanistic analysis to improve the yields and to explore other reactivity possibilities based on this process are currently being studied and will be published in due time.

Supporting Information

Supporting Information File 1

Experimental procedures and analytical data for all substrates and products, product inhibition study, computational calculation information, and relevant energies and Cartesian coordinates.

[<http://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-13-41-S1.pdf>]

Acknowledgements

Funding for this project from the North Carolina Biotechnology Center (BRG-1205) and University of North Carolina at Greensboro is gratefully acknowledged. The authors thank Dr. Franklin J. Moy for assisting with analysis of NMR data and Dr. Daniel A. Todd for acquisition of the high resolution mass spectrometry data at the Triad Mass Spectrometry Laboratory at the University of North Carolina at Greensboro.

References

- Torborg, C.; Beller, M. *Adv. Synth. Catal.* **2009**, *351*, 3027. doi:10.1002/adsc.200900587
- McGlacken, G. P.; Fairlamb, I. J. S. *Eur. J. Org. Chem.* **2009**, 4011. doi:10.1002/ejoc.200900139
- Nicolaou, K. C.; Bulger, P. G.; Sarlah, D. *Angew. Chem., Int. Ed.* **2005**, *44*, 4442. doi:10.1002/anie.200500368
- Terao, J.; Kambe, N. *Acc. Chem. Res.* **2008**, *41*, 1545. doi:10.1021/ar800138a
- Martin, R.; Buchwald, S. L. *Acc. Chem. Res.* **2008**, *41*, 1461. doi:10.1021/ar800036s
- Marion, N.; Nolan, S. P. *Acc. Chem. Res.* **2008**, *41*, 1440. doi:10.1021/ar800020y
- Denmark, S. E.; Regens, C. S. *Acc. Chem. Res.* **2008**, *41*, 1486. doi:10.1021/ar800037p
- Kantchev, E. A. B.; O'Brien, C. J.; Organ, M. G. *Angew. Chem., Int. Ed.* **2007**, *46*, 2768. doi:10.1002/anie.200601663
- Corbet, J.-P.; Mignani, G. *Chem. Rev.* **2006**, *106*, 2651. doi:10.1021/cr0505268
- Schaub, T. A.; Kivala, M. Cross-Coupling Reactions to sp Carbon Atoms. *Metal-Catalyzed Cross-Coupling Reactions and More*; Wiley-VCH Verlag GmbH & Co. KGaA: Weinheim, 2014; pp 665 ff.
- Davies, H. M. L.; Morton, D. *J. Org. Chem.* **2016**, *81*, 343. doi:10.1021/acs.joc.5b02818
- Brückl, T.; Baxter, R. D.; Ishihara, Y.; Baran, P. S. *Acc. Chem. Res.* **2012**, *45*, 826. doi:10.1021/ar200194b
- Stuart, D. R.; Fagnou, K. The Discovery and Development of a Palladium(II)-Catalyzed Oxidative Cross-Coupling of Two Unactivated Arenes. In *Inventing Reactions*; Gooßen, L. J., Ed.; Springer: Berlin, Heidelberg, 2013; pp 91 ff.
- Colby, D. A.; Bergman, R. G.; Ellman, J. A. *Chem. Rev.* **2010**, *110*, 624. doi:10.1021/cr900005n
- Campeau, L.-C.; Rousseaux, S.; Fagnou, K. *J. Am. Chem. Soc.* **2005**, *127*, 18020. doi:10.1021/ja056800x
- Stuart, D. R.; Alsabeh, P.; Kuhn, M.; Fagnou, K. *J. Am. Chem. Soc.* **2010**, *132*, 18326. doi:10.1021/ja1082624
- Gorelsky, S. I.; Lapointe, D.; Fagnou, K. *J. Org. Chem.* **2012**, *77*, 658. doi:10.1021/jo202342q
- Lafrance, M.; Fagnou, K. *J. Am. Chem. Soc.* **2006**, *128*, 16496. doi:10.1021/ja067144j
- Jana, R.; Trivedi, R.; Tunge, J. A. *Org. Lett.* **2009**, *11*, 3434. doi:10.1021/ol901288r
- Tokoroyama, T.; Nakamura, M. *Chem. Lett.* **1977**, *6*, 659. doi:10.1246/cl.1977.659
- Tsuda, T.; Chujo, Y.; Nishi, S.; Tawara, K.; Saegusa, T. *J. Am. Chem. Soc.* **1980**, *102*, 6381. doi:10.1021/ja00540a053
- Tsuda, T.; Okada, M.; Nishi, S.; Saegusa, T. *J. Org. Chem.* **1986**, *51*, 421. doi:10.1021/jo00354a001
- Trost, B. M.; Bunt, R. C. *J. Am. Chem. Soc.* **1998**, *120*, 70. doi:10.1021/ja9726522
- Corey, E. J.; Fraenkel, G. *J. Am. Chem. Soc.* **1953**, *75*, 1168. doi:10.1021/ja01101a047
- Waetzig, S. R.; Rayabarapu, D. K.; Weaver, J. D.; Tunge, J. A. *Angew. Chem., Int. Ed.* **2006**, *45*, 4977. doi:10.1002/anie.200600721
- Waetzig, S. R.; Tunge, J. A. *J. Am. Chem. Soc.* **2007**, *129*, 14860. doi:10.1021/ja077070r
- Grenning, A. J.; Tunge, J. A. *Org. Lett.* **2010**, *12*, 740. doi:10.1021/ol902828p
- Rayabarapu, D. K.; Tunge, J. A. *J. Am. Chem. Soc.* **2005**, *127*, 13510. doi:10.1021/ja0542688
- Sim, S. H.; Park, H.-J.; Lee, S. I.; Chung, Y. K. *Org. Lett.* **2008**, *10*, 433. doi:10.1021/ol702577g
- Pi, S.-F.; Tang, B.-X.; Li, J.-H.; Liu, Y.-L.; Liang, Y. *Org. Lett.* **2009**, *11*, 2309. doi:10.1021/ol900643r
- Torregrosa, R. R. P.; Ariyaratna, Y.; Chattopadhyay, K.; Tunge, J. A. *J. Am. Chem. Soc.* **2010**, *132*, 9280. doi:10.1021/ja1035557

32. Zhang, W.-W.; Zhang, X.-G.; Li, J.-H. *J. Org. Chem.* **2010**, *75*, 5259. doi:10.1021/jo1010284
33. Rodriguez, N.; Goossen, L. J. *Chem. Soc. Rev.* **2011**, *40*, 5030. doi:10.1039/c1cs15093f
34. Gooßen, L. J.; Deng, G.; Levy, L. M. *Science* **2006**, *313*, 662. doi:10.1126/science.1128684
35. Gruber, S.; Zaitsev, A. B.; Wörle, M.; Pregosin, P. S.; Veiros, L. F. *Organometallics* **2008**, *27*, 3796. doi:10.1021/om800295z
36. Yamashita, M.; Hirano, K.; Satoh, T.; Miura, M. *Org. Lett.* **2010**, *12*, 592. doi:10.1021/ol9027896
37. Patel, B. A.; Dickerson, J. E.; Heck, R. F. *J. Org. Chem.* **1978**, *43*, 5018. doi:10.1021/jo00420a029
38. Yamashita, M.; Hirano, K.; Satoh, T.; Miura, M. *Adv. Synth. Catal.* **2011**, *353*, 631. doi:10.1002/adsc.201000897
39. Yamashita, M.; Hirano, K.; Satoh, T.; Miura, M. *Chem. Lett.* **2010**, *39*, 68. doi:10.1246/cl.2010.68
40. Abu Deiab, G. I.; Croatt, M. P. Chapter 3 - Step-Economical Synthesis of Clinprost and Analogs Utilizing a Novel Decarboxylation Reaction. In *Strategies and Tactics in Organic Synthesis*; Michael, H., Ed.; Academic Press, 2017; Vol. 12, pp 95 ff.
41. Nagy, E. E.; Hyatt, I. F. D.; Gettys, K. E.; Yeazell, S. T.; Frempong, S. K., Jr.; Croatt, M. P. *Org. Lett.* **2013**, *15*, 586. doi:10.1021/ol303402e
42. Thadani, A. N.; Rawal, V. H. *Org. Lett.* **2002**, *4*, 4317. doi:10.1021/ol0269594
43. Wilson, S. R.; Zucker, P. A. *J. Org. Chem.* **1988**, *53*, 4682. doi:10.1021/jo00255a007
44. Tang, W.; Prusov, E. V. *Org. Lett.* **2012**, *14*, 4690. doi:10.1021/ol302219x
45. Gagnepain, J.; Moulin, E.; Fürstner, A. *Chem. – Eur. J.* **2011**, *17*, 6964. doi:10.1002/chem.201100178
46. Jeso, V.; Micalizio, G. C. *J. Am. Chem. Soc.* **2010**, *132*, 11422. doi:10.1021/ja104782u
47. Schnermann, M. J.; Romero, F. A.; Hwang, I.; Nakamaru-Ogiso, E.; Yagi, T.; Boger, D. L. *J. Am. Chem. Soc.* **2006**, *128*, 11799. doi:10.1021/ja0632862
48. McCammant, M. S.; Shigeta, T.; Sigman, M. S. *Org. Lett.* **2016**, *18*, 1792. doi:10.1021/acs.orglett.6b00517
49. Todd, D. P.; Thompson, B. B.; Nett, A. J.; Montgomery, J. *J. Am. Chem. Soc.* **2015**, *137*, 12788. doi:10.1021/jacs.5b08448
50. Kong, W.; Che, C.; Kong, L.; Zhu, G. *Tetrahedron Lett.* **2015**, *56*, 2780. doi:10.1016/j.tetlet.2015.04.036
51. Xu, S.; Zhu, S.; Shang, J.; Zhang, J.; Tang, Y.; Dou, J. *J. Org. Chem.* **2014**, *79*, 3696. doi:10.1021/jo500375q
52. Huang, Y.; Fañanás-Mastral, M.; Minnaard, A. J.; Feringa, B. L. *Chem. Commun.* **2013**, *49*, 3309. doi:10.1039/c3cc41021h
53. Schmidt, A.; Hilt, G. *Org. Lett.* **2013**, *15*, 2708. doi:10.1021/ol401015e
54. Lim, H. N.; Parker, K. A. *J. Am. Chem. Soc.* **2011**, *133*, 20149. doi:10.1021/ja209459f
55. Kennard, O.; Watson, D. G.; Fawcett, J. K.; Kerr, K. A. *Tetrahedron* **1970**, *26*, 607. doi:10.1016/S0040-4020(01)97853-6
56. Hettrick, C. M.; Kling, J. K.; Scott, W. J. *J. Org. Chem.* **1991**, *56*, 1489. doi:10.1021/jo00004a028
57. Gradén, H.; Hallberg, J.; Kann, N.; Olsson, T. *J. Comb. Chem.* **2004**, *6*, 783. doi:10.1021/cc049929w
58. Surovtseva, D. A.; Orlov, D. A.; Morozova, T. A.; Krylov, A. V.; Belov, A. P. *Kinet. Catal.* **2006**, *47*, 855. doi:10.1134/s0023158406060073
59. Alfonso, D. R.; Jordan, K. D. *J. Comput. Chem.* **2003**, *24*, 990. doi:10.1002/jcc.10233

License and Terms

This is an Open Access article under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

The license is subject to the *Beilstein Journal of Organic Chemistry* terms and conditions: (<http://www.beilstein-journals.org/bjoc>)

The definitive version of this article is the electronic one which can be found at: [doi:10.3762/bjoc.13.41](https://doi.org/10.3762/bjoc.13.41)