RSC Advances



PAPER

View Article Online



Cite this: RSC Adv., 2024, 14, 38285

π - π stacking assisted regioselectivity regulation in palladium-catalyzed cyclization reactions: a theoretical study†

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The regulation of regioselectivity is an objective often pursued by organic chemists, and the comprehension of its mechanisms is crucial for devising efficient synthetic pathways. In this report, we conducted theoretical calculations to explore the regioselectivity regulatory mechanisms of two palladiumcatalyzed cyclization reactions. In these cyclization reactions, manipulating the structural differences in the reaction substrates leads to the formation of distinct products. A detailed reaction mechanism and reactivity profile for this reaction were revealed. Furthermore, a $\pi - \pi$ stacking assisted regionselectivity regulatory mechanism was proven by distortion-interaction energy analysis and noncovalent interaction calculations. The calculated results presented herein provide a theoretical guide for further experimental investigations of regioselectivity regulation.

Received 11th September 2024 Accepted 27th November 2024

DOI: 10.1039/d4ra06552b

rsc li/rsc-advances

Introduction

Regioselectivity is an important field in organic synthetic chemistry that mainly focuses on the differences in reactivity at various positions on a substrate.1 Several reactive sites may exhibit similar activity, leading to the formation of a mixture of isomers in the final product. Catalytic methods are ideal synthetic approaches in organic chemistry, as they allow for the prediction switching of regioselectivity to access site-diverse regioisomers by altering the fewest reaction parameters.2

Various regulatory strategies in organic chemistry, such as steric effects,3 electronic effects,4 chelation control,5 and ligand control,6 have been used to achieve regioselectivity7 (Scheme 1a). Chemists can use these strategies to manipulate the outcome of organic reactions to achieve certain modifications at specific sites within complex molecules, but the regulatory mechanism of regioselectivity is often ambiguous. In organic synthesis, understanding the regulatory mechanisms of these strategies is essential for designing efficient synthetic routes to obtain the targeted products.8

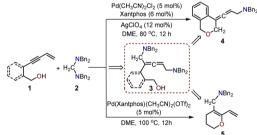
Recently, Huang et al. described two palladium (Pd)catalyzed cyclization reactions of enynols with aminals.9 In their reports, different products were obtained by manipulating the skeletal variances of substrates. As shown in Scheme 1b, the

use of (2-(but-3-en-1-yn-1-yl)phenyl)methanol as the substrate

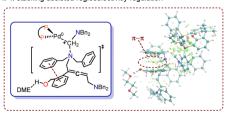
led to the predominant formation of the O-heterocyclecontaining allenic amine product 4. Furthermore, the use of

hept-6-en-4-yn-1-ol as the substrate resulted in the formation of

polysubstituted 1,3-diene embedded in O-heterocycle 5 as the main product (Scheme 1b). Control experiments confirmed the



(c) π - π stacking assisted regional regulation



Scheme 1 Palladium-catalyzed cyclization of enynols with aminals.

non-cyclic allenic 1,5-diamine 3 as the key intermediate in both cyclization reactions. However, the detailed regioselectivity (a) Regulatory strategies for regioselectivity Steric effects **Electronic effects Chelation control** Conformational control (b) Experiment observations

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[†] Electronic supplementary information (ESI) available. DOI: https://doi.org/10.1039/d4ra06552b

regulatory mechanism for these Pd-catalyzed cyclizations is still not clear. Herein, we used density functional theory (DFT) calculations to explore these Pd-catalyzed cyclization reactions and proposed a π - π stacking assisted regioselectivity regulatory mechanism (Scheme 1c). A detailed reaction mechanism and reactivity profile were also provided.

Computational methods

All DFT calculations were carried out with Gaussian 16.10 The B3LYP functional11 with a standard 6-31G(d) basis set (the SDD basis set for Pd) was used for geometry optimizations. Harmonic frequency calculations were performed for all stationary points to confirm them as local minima or transition structures and to derive the thermochemical corrections of the enthalpies and free energies. All minima had zero imaginary frequency and all transition states had only one imaginary frequency. The solvent effects were considered by single-point calculations on the gas-phase stationary points with the SMD continuum solvation model.12 The M06 (ref. 13) functional with the 6-311+g(d) basis set (the SDD basis set for Pd) was used to calculate the single-point energies with dimethyl ether as the solvent. For the 1,2-dimethoxyethane (DME) solvent data not included in Gaussian 16, the key parameters were set in the input file manually (for 1,2-dimethoxyethane, eps = 7.2, while the other parameters were the same as those for diethyl ether). The energies given in this report were the M06 calculated Gibbs free energies in 1,2-dimethoxyethane solvent. The optimized structures were displayed using CYLview.14 The distortion and interaction energies were calculated B3LYP functional with the 6-31G(d) basis set. Additionally, a basis set superposition error (BSSE)¹⁵ correction was applied when calculating the interaction energies. The noncovalent interactions (NCIs)16 were calculated using B3LYP-D3 functional with the 6-31G(d) basis set.

The global reactivity index values were also calculated using the M06 functional and the 6-311+g(d) basis set for all atoms. The N (global nucleophilicity index)¹⁷ values were calculated with eqn (1), where $\varepsilon_{\text{HOMO}}$ refers to the orbital energy of the highest occupied molecular orbital (HOMO) and $\varepsilon_{\text{LOMO}}$ refers to the orbital energy of the lowest unoccupied molecular orbital (LUMO). In eqn (1), tetracyanoethylene (TCE) with the lowest HOMO energy among the organic molecules was used as a ref. 18.

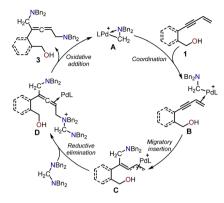
$$N = \varepsilon_{\text{HOMO}} (\text{Nu}) - \varepsilon_{\text{HOMO}} (\text{TCE})$$
 (1)

The $N_{\rm k}$ (local nucleophilicity index) values were calculated according to eqn (2), where the $f_{\rm k}^-$ (condensed Fukui function) values can be obtained by $f_{\rm k}^- = q_{\rm k}(N-1) - q_{\rm k}(N)$.¹⁹

$$N_{\mathbf{k}} = N \times f_{\mathbf{k}}^{-} \tag{2}$$

Results and discussion

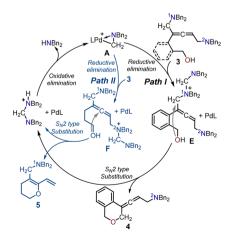
The proposed catalytic cycle for the Pd-catalyzed cyclization of enynols with aminals is illustrated in Schemes 2 and 3. Scheme



Scheme 2 The proposed catalytic cycle for the production of intermediate 3.

2 depicts the catalytic cycle for the allenic 1,5-diamine intermediate 3. The aminomethyl cyclopalladated complex **A** was the active catalyst in both cyclizations. The coordination of the terminal alkene in the enynol substrate **1** with the Pd(π) center generated intermediate **B**. The subsequent migratory insertion of the enyne triple bond into the Pd(π)-C bond provided the π -allylpalladium intermediate **C**, which was intercepted by an aminal to form intermediate **D** νia an S_N2-type reductive elimination process. Finally, the S_N2-type oxidative addition of the amino cation to the Pd(0) center afforded the non-cyclic allenic 1,5-diamine intermediate **3** together and regenerated the active Pd complex **A** to complete the catalytic cycle.

The proposed catalytic cycles in the production of the allenic amine 4 and the 1,3-diene O-heterocycle product 5 are depicted in Scheme 3. By employing (2-(but-3-en-1-yn-1-yl)phenyl) methanol 1 as the substrate in Path I, the 1N atom in intermediate 3 can serve as a nucleophilic site to interact with the cyclopalladated complex A through S_N2 -type reductive elimination, resulting in the formation of the quaternary ammonium intermediate E and Pd(0). The intermediate E is intramolecularly intercepted by the pendent alcohol through S_N2 -type substitution to yield the desired product 4. Finally, the S_N2 -type



Scheme 3 Proposed catalytic cycle for the generation of the allenic amine product 4 and 1,3-diene *O*-heterocycle 5.

oxidative addition of the generated amino cation to the Pd(0) center regenerated the active palladium-complex A to complete the catalytic cycle. Moreover, the use of hept-6-en-4-yn-1-ol 1' as the substrate led to the 2N atom in intermediate 3 acting as a nucleophilic site to attack the Pd complex A, forming the quaternary ammonium intermediate F. The allene in intermediate F was intercepted by the pendent alcohol to give the 1,3diene O-heterocycle product 5.

The calculated free energy profiles for the generation of intermediate 3 using (2-(but-3-en-1-yn-1-yl)phenyl)methanol 1 as the substrate are presented in Fig. 1, with the aminomethyl cyclopalladated complex INT1 serving as the reference zero point. The binding of the terminal alkene in the enynol substrate 1 to the Pd(II) center led to the endergonic generation of intermediate INT2 (12.9 kcal mol^{-1}). The subsequent migratory insertion of the enyne triple bond into the Pd(II)-C bond took place through the transition state TS1, resulting in the formation of the π -allylpalladium intermediate INT3, with a free energy barrier of 12.2 kcal mol⁻¹. In transition state **TS1**, the length of the formed C-C bond is 2.19 Å. The intermediate INT3 would then be intercepted by an aminal, leading to the formation of allene-coordinated Pd(0) intermediate INT4 through an S_N2-type reductive elimination transition state TS2 (activation free energy = $27.0 \text{ kcal mol}^{-1}$). In the transition state TS2, the length of the formed C-N bond is 1.97 Å. The following S_N2-type oxidative addition occurred between Pd(0) and the amino cation via transition state TS3, with an energy barrier of 4.4 kcal mol⁻¹. The release of the allenic 1,5-diamine intermediate 3 led to the exothermic formation of the aminomethyl cyclopalladated complex INT1 (4.3 kcal mol⁻¹) in comparison with substrates 1 and 2. Moreover, the calculated free energy profiles for the generation of intermediate 3' using hept-6-en-4yn-1-ol 1' as the substrate are presented in Fig. S1.†

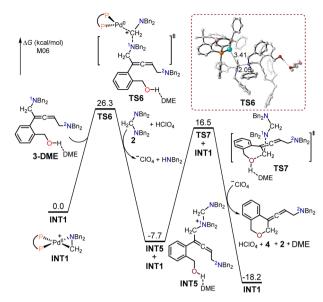


Fig. 2 Calculated free energy profiles for the production of the allenic amine product 4 using (2-(but-3-en-1-yn-1-yl)phenyl)methanol 1 as the substrate.

The π -allylpalladium in INT3 can be intercepted by the tethered alcohol to form the 1,3-diene O-heterocycle product 5 through the S_N2-type reductive elimination transition states TS4 and TS5. In TS4, the aminal substrate functioned as a base to facilitate reductive elimination, while the intramolecular amido group in transition state TS5 assisted in reductive elimination. The energies of TS4 and TS5 were respectively 1.1 and 21.6 kcal mol⁻¹ higher than that of **TS2**, indicating that these pathways were unfavorable.

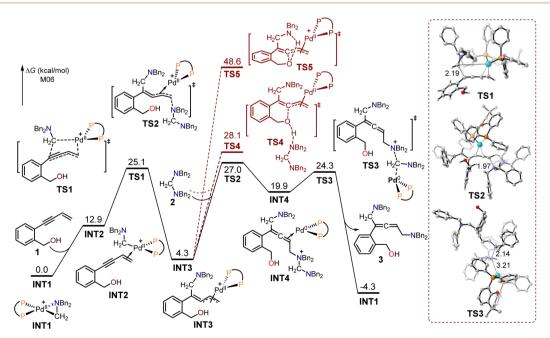


Fig. 1 Calculated free energy profiles for the production of intermediate 3 using (2-(but-3-en-1-yn-1-yl)phenyl)methanol 1 as the substrate.

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Paper

The free energy profiles for the generation of the allenic amine product 4, using (2-(but-3-en-1-vn-1-vl)phenyl)methanol 1 as the substrate are presented in Fig. 2. To investigate the effect of solvation on the system's structure and energy, an explicit DME molecule was included with substrate 3 to allow for the possibility of intermolecular hydrogen bonding between the hydroxyl group of 3 and a solvent oxygen atom. In the absence of the solvent molecule, an artefactual and misleading intramolecular O-H··· π hydrogen bond would occur. The ¹N atom in intermediate 3-DME served as a nucleophilic site to interact with the cyclopalladated complex INT1 through S_N2type reductive elimination to provide the quaternary ammonium intermediate INT5 and Pd(0). The free energy barrier for this S_N2-type reductive elimination transition state TS6 is 26.3 kcal mol⁻¹; the formed C-N bond and broken C-Pd bond had lengths of 2.05 and 3.41 Å, respectively. The resulting Pd(0) was captured by the amino cation, produced through the protonation of the aminal with HClO4, to form the cyclopalladated complex, INT1. The generated intermediate INT5 was then intramolecularly intercepted by the pendent alcohol through the S_N2-type substitution transition state TS7 with a free energy barrier of 24.2 kcal mol⁻¹, to yield the desired allenic amine product 4.

The free energy profiles for the formation of the 1,3-diene Oheterocycle product 5' using (2-(but-3-en-1-yn-1-yl)phenyl) methanol 1 as the substrate are depicted in Fig. 3. The ²N atom in intermediate 3-DME acted as a nucleophilic site to undergo S_N2-type reductive elimination with the cyclopalladated complex INT1 to generate the quaternary ammonium intermediate INT6. The free energy barrier for the S_N2type reductive elimination transition state TS8 (28.8 kcal mol⁻¹) was 2.5 kcal mol⁻¹ higher than that of **TS6**, as the use of (2-(but-3-en-1-yn-1-yl)phenyl)methanol as the substrate primary resulted in the formation of the allenic amine product 4. In TS8, the

formed C-N bond had a length of 2.13 Å, while the broken C-Pd bond was 3.26 Å. The intermediate **INT6** is expected to undergo an intramolecular interception by the pendant alcohol through an S_N 2-type substitution transition state **TS9**, resulting in the formation of the 1,3-diene O-heterocycle product 5'.

The free energy profiles for the formation of the allenic amine product 4' using hept-6-en-4-yn-1-ol 1' as the substrate are depicted in Fig. 4. The ¹N atom in intermediate 3'-DME functioned as a nucleophilic site that interacted with the cyclopalladated complex INT1 via S_N2-type reductive elimination to create the quaternary ammonium intermediate INT5' and Pd(0). The energy barrier for this S_N2-type reductive elimination transition state TS6' was 33.1 kcal mol⁻¹. The resulting intermediate INT5' was further reacted intramolecularly with the adjacent alcohol via an S_N 2-type substitution transition state TS7', with a free energy barrier of 27.7 kcal mol⁻¹, to yield the allenic amine product $\mathbf{4}'$.

The free energy profiles for the synthesis of the 1,3-diene Oheterocycle product 5 from the hept-6-en-4-yn-1-ol 1' substrate are displayed in Fig. 5. The nitrogen atom (2N) in intermediate 3'-DME functioned as a nucleophilic site to undergo S_N2-type reductive elimination with the cyclopalladated complex INT1, producing the quaternary ammonium intermediate INT6'. The energy barrier for the S_N2-type reductive elimination transition state TS8' was 31.8 kcal mol⁻¹. The intermediate INT6' underwent intramolecular interception by the adjacent alcohol through an S_N2-type substitution transition state TS9', with a free energy barrier of 29.9 kcal mol⁻¹, to yield the 1,3-diene Oheterocycle product 5. TS8' had a free energy 1.3 kcal mol⁻¹ lower than that of TS6', and the free energy of the 1,3-diene Oheterocycle product 5 was 12.4 kcal mol⁻¹ lower than that of the allenic amine product 4'. These findings elucidate that the primary product formed was the 1,3-diene O-heterocycle product 5 when using hept-6-en-4-yn-1-ol as the substrate.

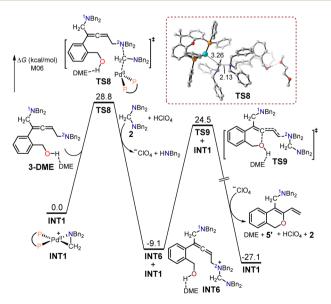


Fig. 3 Calculated free energy profiles for the production of the 1,3diene O-heterocycle product 5' using (2-(but-3-en-1-yn-1-yl)phenyl) methanol 1 as the substrate

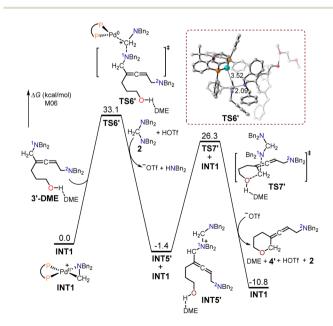


Fig. 4 Calculated free energy profiles for the production of the allenic amine product 4' using hept-6-en-4-yn-1-ol 1' as the substrate.

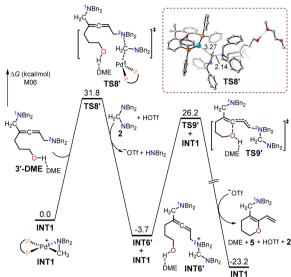
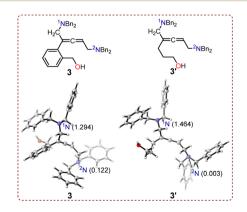


Fig. 5 Calculated free energy profiles for the production of the 1,3diene O-heterocycle product 5 using hept-6-en-4-yn-1-ol 1' as the substrate.

The free energy barrier for **TS6** was 6.8 kcal mol⁻¹ lower than that of TS6', whereas the free energy barriers for TS8 was 3.0 kcal mol⁻¹ lower than that of TS8'. The reactivity of the ¹N atom decreased when the substrate was changed from (2-(but-3en-1-yn-1-yl)phenyl)methanol 1 to hept-6-en-4-yn-1-ol 1', whereas the reactivity of the ²N atom remained unchanged. This difference in the reactivities of ¹N and ²N contributed to the varying selectivity observed with different substrates.

The local nucleophilicity index N_k values were calculated to analyze the nucleophilicity of the nitrogen atoms (¹N and ²N) in intermediates 3 and 3'. The calculated N_k values for 1N and 2N in intermediate 3 were respectively 1.294 and 0.122 eV, while the values were 1.464 and 0.003 eV for intermediate 3' (Fig. 6). The nucleophilicity of ¹N was stronger than that of ²N in both intermediates. Notably, the nucleophilicity of ¹N and ²N in noncyclic allenic 1,5-diamine intermediates 3 and 3' remained unaffected when using (2-(but-3-en-1-yn-1-yl)phenyl)methanol 1 or hept-6-en-4-yn-1-ol 1' as substrates.



Calculated local nucleophilicity index N_k values.

Table 1 Distortion and interaction energies of the transition states **TS6** and **TS6**′. Values are given in kcal mol $^{-1}$ ($\Delta E_{\rm dist}^{\uparrow}$ is the distortion energies of the transition states **TS6**′. gies. $\Delta E_{\text{int}}^{\ddagger}$ is the interaction energies. ΔE^{\ddagger} is the reaction energies)

	$\Delta E_{ m dist(3-DME}^{\ddag}$ or 3'-DME)	$\Delta \mathit{E}_{\mathrm{dist}(\mathbf{INT1})}^{\ddagger}$	$\Delta E_{\rm dist}^{\ddagger}$	$\Delta E_{ m int}^{\ddagger}$	ΔE^{\ddagger}
TS6	9.5	35.9	45.4	-15.9	29.5
TS6'	11.2	37.2	48.4	-16.5	32.0

Distortion-interaction energy analysis was used to elucidate the disparity in the free energy barrier between transition states **TS6** and **TS6**'. The distortion energy of **TS6**' was 3.0 kcal mol⁻¹ higher than that of TS6, while their interaction energies were similar (Table 1). The distortion energies of intermediates 3-**DME** and 3'-**DME** in **TS6** and **TS6'** were 9.5 and 11.2 kcal mol $^{-1}$. respectively. The distortion energy of INT1 in both TS6 and TS6' were similar. Analysis of the distortion energy revealed that the difference in energy barriers of TS6 and TS6' was controlled by the distortion energies of intermediates 3-DME and 3'-DME.

The structural information for intermediate 3-DME and transition state TS6 are shown in Fig. 7. In the optimized structure of intermediate 3-DME, the dihedral angle of ¹C-¹N-²C-³C is 95.7°, whereas this dihedral angle changes to 54.7° in **TS6**. In the optimized structure of intermediate 3′-**DME**, the dihedral angle of ${}^{1}C^{-1}N^{-2}C^{-3}C$ is -64.8° , which changes to 55.2° in TS6' (Fig. 8). These calculated results indicated that the difference in free energy barrier between TS6 and TS6' was controlled by the distortion of the benzyl group associated with ¹N in intermediates 3-DME and 3'-DME.

To further investigate the regioselectivity regulatory mechanism behind these Pd-catalyzed cyclization reactions, NCI analysis was conducted for transition states TS6 and TS6'. The π - π stacking of two phenyl rings in transition state **TS6** occurred when (2-(but-3-en-1-yn-1-yl)phenyl)methanol 1 was used as the substrate (Fig. 9). By contrast, in transition state **TS6'**, there is solely a C-H··· π interaction between the alkyl chain and phenyl group. NCI calculations revealed that the π - π stacking in TS6 are stronger than the C-H $\cdots\pi$ interactions in

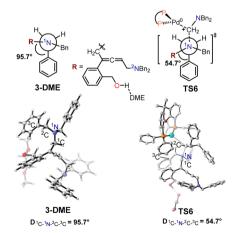


Fig. 7 Structural information for intermediate 3-DME and transition state TS6

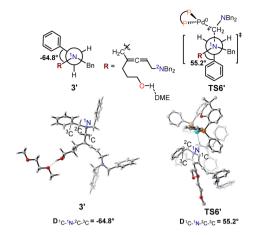


Fig. 8 Structural information for intermediate 3'-DME and transition state TS6'.

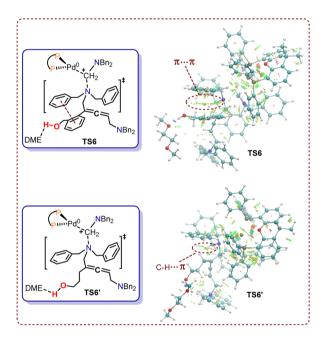


Fig. 9 Noncovalent interactions of TS6 and TS6' (blue, attraction; green, weak interaction; red, steric effect).

TS6′. This difference results in a lower distortion energy for **TS6** compared to **TS6**′, which consequently reduces the reactivity of the ¹N atom when the substrate is changed from (2-(but-3-en-1-yn-1-yl)phenyl)methanol 1 to hept-6-en-4-yn-1-ol 1′.

Conclusions

In this work, DFT calculations were used to explore the mechanism of Huang's Pd-catalyzed cyclization reactions with an emphasis on the regioselectivity regulation mechanism. Theoretical calculations revealed that these reactions proceeded via migratory insertion, S_N 2-type reductive elimination, and oxidative addition to give the non-cyclic allenic 1,5-diamine intermediate. The allenic amine and 1,3-diene O-heterocycle

products were then obtained through subsequent reductive elimination, S_N 2-type substitution, and oxidative addition.

The reactivity of the 1N atom decreased when (2-(but-3-en-1-yn-1-yl)phenyl)methanol instead of hept-6-en-4-yn-1-ol was used as the substrate, while the reactivity of the 2N atom exhibited minimal change. This difference in reactivities of 1N and 2N contributed to the varying selectivity observed with different substrates. Distortion–interaction energy analysis and NCI calculations indicated that the presence of π – π stacking in intermediate 3-DME led to the lower distortion energy of TS6 than that of TS6', consequently reducing the reactivity of the 1N atom upon changing the substrate from (2-(but-3-en-1-yn-1-yl) phenyl)methanol to hept-6-en-4-yn-1-ol for regioselective control.

Data availability

The data that support the findings of this study have been included in the main text and ESI† and are available from the corresponding author upon reasonable request.

Author contributions

S. Liu conceived and designed the project. S. Liu designed the computational studies. D. Zhang, Y. Gong, L. Ma and L. Li performed the DFT calculations. S. Liu and W. Chen prepared the manuscript, D. Zhang, Y. Gong, L. Ma and L. Li prepared the ESI.†

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (No. 22303010 and 22271034). We are thankful to the Scientific Research Fund of Chongqing University of Arts and Sciences (no. R2020SHH01) for financial support.

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