Mechanistic Insight Into the AuCN Catalyzed Annulation Reaction of Salicylaldehyde and Aryl Acetylene: Cyanide Ion Promoted Umpolung Hydroacylation/Intramolecular Oxa-Michael Addition Mechanism

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The detailed mechanism of the AuCN-catalyzed annulation of salicylaldehyde (SA) and phenyl acetylene leading to isoflavanone-type complexes has been investigated via density functional theory (DFT) calculations. Reaction pathways and possible stationary points are obtained with the combined molecular dynamics and coordinate driving (MD/CD) method. Our calculations reveal that the cyanide ion promoted umpolung hydroacylation/intramolecular oxa-Michael addition mechanism is more favorable than the Au(I)/Au(III) redox mechanism proposed previously. In the umpolung mechanism, the hydroxyl of SA is found to strongly stabilize the cyanide ion involved intermediates and transition states via hydrogen bond interactions, while the Au(I) ion always acts as a counter cation. The overall reaction is exergonic by 41.8 kcal/mol. The hydroacylation of phenyl acetylene is the rate-determining step and responsible for the regioselectivity with a free energy barrier of 27.3 kcal/mol. These calculated results are in qualitative accord with the experimental findings.

Keywords: DFT, MD/CD, cyanide ion, umpolung, hydroacylation

INTRODUCTION

The catalytic annulation of chelating aldehydes and unsaturated hydrocarbons provides a general, highly efficient, and atom-economical strategy to synthesize complex carbocyclic and heterocyclic frameworks (Willis, 2010). Salicylaldehyde (SA) is a highly favored chelating aldehyde because of its ready availability as well as common ortho-hydroxy or alkoxy arylcarbonyl motifs in natural products and bioactive molecules. In recent years, starting from SA, various transition-metal-catalyzed Csp2-H activation and heterocyclization reactions have been developed to efficiently construct various important heterocycles of biologically active compounds and drug molecules (Shimizu et al., 2008; Zeng and Li, 2014; Baruah et al., 2016; Yang and Yoshikai, 2016). For example, the Rh(III)-catalyzed (Shimizu et al., 2008), Ru(II)-catalyzed (Baruah et al., 2016), Co(I)-diphosphine-catalyzed (Yang and Yoshikai, 2016) Csp2-H activation and annulation reactions of SA with monosubstituted...
and disubstituted alkyne have been reported for the synthesis of chromone derivatives.

Over the last two decades, homogeneous gold catalysis has developed very rapidly and broadly in organic synthesis due to its rich chemistry and fascinating reactivity. It has emerged as a very efficient method for rapid construction of complex and highly functionalized molecules (Fukuda and Utimoto, 1991; Teles et al., 1998; Hashmi et al., 2000a,b; Yao and Li, 2004; Nguyen et al., 2006; Boorman and Larrosa, 2011; Xie et al., 2014; Dorel and Echavarren, 2015; Joost et al., 2015; Qian and Zhang, 2015; Kumar and Nevado, 2017; Akram et al., 2018; Mandal and Datta, 2018; Matsumoto et al., 2018; Wu et al., 2018; Kreuzahler et al., 2019; Mascareñas et al., 2019). Very recently, starting with SA and aryl acetylene, Li and co-workers developed an AuCN-catalyzed system (Skouta and Li, 2007) to afford the isoflavanone-type frameworks, which have many possible applications in the synthesis of isoflavanone natural products. Although the AuCN-catalyzed annulation of SA and aryl acetylene can effectively construct isoflavanone-type frameworks, the reaction conditions are very harsh. It takes place at high temperature (150 °C), and over-stoichiometric amounts of alkynes (3 equivalent) are required, as shown in Scheme 1A. Moreover, the detailed mechanism of this reaction is still elusive.

We believe that the elucidation of the mechanism of this reaction is helpful for designing new and milder catalytic reaction systems. Based on the regioselectivity of the AuCN-catalyzed annulation reaction, Li and co-workers suggested an Au(I)/Au(III) redox mechanism via Csp2-H activation of aldehyde, as shown in Scheme 1B. This mechanism starts with an oxidative addition of the aldehyde Csp2-H bond leading to an acyl Au(III) hydride I. Then a hydrometalation step occurs, followed by a reductive elimination process and then an intramolecular conjugate addition, affording the final isoflavanone derivative and regenerating the Au(I) catalyst. Experiments indicated that the phosphine ligands had significant influence on the yield of product. As shown in Scheme 1A, triethylphosphane (PET₃) and tributylphosphane (PBu₃) exhibited moderate to good efficiency (36 and 78% yield, respectively), while other alternative ligands, such as triphenylphosphane (PPh₃) and tri-tert-butylphosphane (P-t-Bu₃), providing only trace amounts of product.

In the present work, we carry out density functional theory (DFT) calculations to investigate the mechanism of the AuCN-catalyzed annulation reaction between SA and phenyl acetylene using AuCN as the catalyst and PET₃ as the model ligand. The combined molecular dynamics and coordinate driving (MD/CD)
method (Yang et al., 2017b, 2018) developed by us is applied to explore reaction pathways and determine the structures of all possible stationary points along reaction pathways. This method is a cost-effective theoretical tool for automatically searching low-energy reaction pathways for relatively large chemical reactions in both gas phase and solution. Our calculations indicate that the pathway initiated by the oxidative addition of aldehyde Csp2-H bond toward Au (I) center could be excluded, due to its high activation barrier. Instead, our calculations suggest a cyanide ion promoted umpolung hydroacetylation/intramolecular oxa-Michael addition mechanism for this reaction.

**RESULTS AND DISCUSSIONS**

**Au(I)/Au(III) Redox Mechanism**

Initially, we investigated the “intuitive” Au(I)/Au(III) redox mechanism, which involves the direct oxidative addition of aldehyde Csp2-H bond to PEt3AuCN complex 1. As shown in Figure 1, the formation of a hydrogen bond between cyano group of PEt3AuCN and the hydroxyl of SA in the complex 5 is endergonic by 0.9 kcal/mol. However, the direct oxidative addition of the formyl Csp2-H bond of SA toward the Au(I) center via transition state TS5/Ia to form the Au(III) intermediate Ia is required to overcome a high activation barrier of 44.4 kcal/mol (relative to reactants), which is inconsistent with the experimental conditions that the reaction occurs at 150°C. Therefore, instead of Au(I)/Au(III) redox catalytic cycle proposed previously, there might be an energetically more favorable pathway.

**The Cyanide Ion Promoted Umpolung Pathway**

**Determination of the Active Catalyst**

According to the previous experimental studies (Hormann et al., 1986), AuCN can readily react with a molecular phosphine ligand to generate a linear PEt3AuCN species 1. It is generally accepted that linear Au(I) could not associate with a second or third ligand because this is an entropy decreasing process (Schwerdtfeger et al., 2003; Carvajal et al., 2004). However, our calculations show that the association of PEt3AuCN with two additional PEt3 ligands to form a tetracoordinated Au(I) species 3 is exergonic by 9.9 kcal/mol (relative to complex 1, see Scheme 2). This result indicates the formation of the tetracoordinated Au (I) highly exergonic, free energies discussed in this work are with respect to PEt3AuCN.
complex 3 is possible. In fact, in the related experimental studies (Skouta and Li, 2007), 25-fold amounts of phosphine ligand (with respect to AuCN) is required to achieve a good yield of the isoflavanone derivatives. These theoretical and experimental results suggest that the tetracoordinated Au (I) complex 3 could be formed under the experimental conditions. Although the
reaction of Au (I) complex 3 with another PEt$_3$ molecule could generate a zwitterionic intermediate (PEt$_3$)$_2$PAu$^+...$CN$^-$, this process is thermodynamically unfavorable (highly endergonic by 15.1 kcal/mol). Thus, we assume that the tetracoordinated complex 3 might be involved in the catalytic cycle. The elongation of Au-CN bond length in (PEt$_3$)$_2$PAuCN by 0.14 Å (compared to PEt$_3$AuCN) indicates the Au-CN bond strength in (PEt$_3$)$_2$PAuCN is weakened, increasing the nucleophilic reactivity of cyanide ion.

HCN Promoted Formation of $\alpha$-Cyano Carbanion Intermediate

With (PEt$_3$)$_3$AuCN as the active catalyst, we found that the cyano group of catalyst 3 could react with the phenolic hydroxyl group of SA to liberate the HCN molecule, which then occurs nucleophilic addition toward the carbonyl of SA to generate a $\alpha$-cyano carbanion intermediate 13. The Gibbs free energy profiles for these processes are presented in Figures 2, 3, respectively. The optimized geometries of the involved species along these reaction pathways are displayed in Figure 4.

First, complexation of the catalyst 3 and SA forms the complex 6, which is stabilized by a O-H...N hydrogen bond interaction ($d_{O-H...N} = 1.80$ Å) and exergonic by 0.4 kcal/mol, as shown in Figure 2. Then, the dissociation of cyanide ion from the Au(I) center in complex 6 leads to an [Au(I)]$^+...$CN$^-$ ion-pair complex 7 (via the transition state TS$_{6/7}$). Both the transition state TS$_{6/7}$ and ion-pair complex 7 could be stabilized by strong hydrogen bond interaction with the phenolic hydroxyl of SA, in which the distances of O-H...CN$^-$ are 1.79 and 1.63 Å, respectively. This process is slightly endergonic by 7.0 kcal/mol with a barrier of 9.3 kcal/mol. Subsequently, complex

![Figure 4](https://example.com/figure4.png) Optimized geometries of some species involved in the formation of the HCN and $\alpha$-cyano carbanion intermediate 13. Hydrogen atoms except for those involved in the reaction are omitted for clarity. All bond lengths are in Å.
7 undergoes a proton transfer process from the acidic phenolic hydroxyl to the cyanide ion to generate a new ternary ion-pair complex 8, HCN-SA−-(PET$_3$)$_3$Au$^+$. The liberation of HCN from the complex 8 forms the complex 9, SA−-(PET$_3$)$_3$Au$^+$. Overall, the formation of the separated HCN and complex 9 is endergonic by 9.3 kcal/mol with a free energy barrier of only 9.8 kcal/mol.

**FIGURE 5** | Gibbs free energy profile for the hydroacylation of phenylacetylene process. Relative free energies (with respect to separated reactants) are given in kcal/mol.

**FIGURE 6** | Optimized geometries of some species involved in the hydroacylation of phenylacetylene. All bond lengths are in Å.
kcal/mol (with respect to complex 6). These results indicate that a free HCN molecule may be generated under the experimental conditions (150 °C).

For the species 7, it may convert into a new complex 7’, in which the cationic gold center of (PEt$_3$)$_3$Au$^+$ is associated with the oxygen atom of carbonyl group ($d_{\text{Au-O}} = 2.86$ Å) (see Figure 3). Cyanide ion in complex 7’ subsequently attacks the carbonyl carbon atom of SA (with the assistance of the phenolic hydroxy), leading to a gold-ligated alkoxide intermediate 10 via the transition state TS$_{7'/10}$ (with a barrier of 25.4 kcal/mol). Then, the association of the basic alkoxide intermediate 10 and a free HCN (generated through the pathway described above) forms a hydrogen bond stabilized complex 11 ($d_{\text{O-H-CN}} = 1.39$ Å). The following intramolecular proton transfer leads...

![Figure 7](image1.png)

**FIGURE 7** | Gibbs free energy profile of the intramolecular oxa-Michael process in toluene solution. Relative free energies (with respect to separated reactants) are given in kcal/mol.

![Figure 8](image2.png)

**FIGURE 8** | Optimized geometries of some species involved in the intramolecular oxa-Michael process. All bond lengths are in Å.
to an Au$^+$-cyanohydrin-CN$^-$ complex 12 (via transition state TS$_{11/12}$). The cyanide ion in complex 12, stabilized by hydrogen bond interaction with cyanohydrin, could abstract the α-H of cyanohydrin to form α-cyano carbanion intermediate 13 and regenerate the HCN simultaneously. The free energy barrier for the CN$^-$ assisted proton abstraction (via transition state T$_{212/13}$) is 24.7 kcal/mol. In addition, the pathway involving SA-catalyzed intramolecular 1,2 H-shift of the gold-ligated alkoxide intermediate 10 could also form α-cyano carbanion intermediate 13 (via transition state TS$_{12/13}$−SA, see Figure S1 for details). In summary, the HCN promoted carbonyl umpolung process is endergonic by 24.7 kcal/mol (relative to intermediate 6). The rate-limiting step for this stage is the nucleophilic attack of cyanide ion at the carbonyl carbon atom with an activation barrier of 25.4 kcal/mol.

Other reaction channels to form the alkoxide intermediate 10, starting from complex 6 or 9, could also be located with the MD/CD method (see Figure S2 for details). However, the related free energy barriers for these pathways are higher by 3−5 kcal/mol than the pathway listed in Figure 3. Therefore, the pathway, starting from complexes 7 or 7′ is responsible for the formation of alkoxide intermediate 10, which then converts into α-cyano carbanion intermediate 13.

**Hydroacylation of Phenylacetylene**

Addition of cyanide ion to SA leads to an umpolung of the carbonyl SA, and the corresponding α-cyano carbanion intermediate 13 could act as an acyl anion equivalent to react with phenylacetylene 14, forming a branched α-β unsaturated ketone complex 17 and regenerating the cyanide ion. The Gibbs free energy profile for this process is presented in Figure 5. The optimized geometries of some stationary points are displayed in Figure 6.

First, complexation of the phenyl acetylene 14 with the α-cyano carbanion intermediate 13 forms an intermolecular complex 15. Then, a branched hydroacylation intermediate 16-bra could be generated via a concerted alkyne-Ene transition state (Piel et al., 2011; Schedler et al., 2013) (transition state TS$_{15/16-bra}$), in which, the proton transfer from the α-cyano carbanion intermediate to the terminal carbon (C$_1$) of phenylacetylene is accomplished with a nucleophilic attack of the acetyl anion center at the C$_2$ position of phenyl acetylene. The formation of intermediate 16-bra is exergonic by 33.2 kcal/mol with a barrier of 27.3 kcal/mol. Subsequently, the elimination of the cyanide ion from intermediate 16, with the assistance of the phenolic hydroxyl (via transition state TS$_{16-bra/17}$), forms a branched α-β unsaturated ketone complex 17 (Other possible reaction pathways starting from intermediate 16-bra are presented in Figure S3). This ternary intermediate 17 was stabilized by multiple non-covalent interactions, including the electrostatic interaction between the Au cation and cyanide ion ($d$$_{Au-CN}$ = 3.12 Å) and a strong hydrogen bond interaction ($d$$_{O−H...CN}$ = 1.65 Å) formed between the cyanide ion and phenolic hydroxyl group. Along the energy profile described...
above, we can find that this hydroacylation process is exergonic by 45.7 kcal/mol, relative to the intermediates 13 and 14. The concerted alkyne-Ene process is the rate-determining step with a free energy barrier of 27.3 kcal/mol.

Intramolecular Oxa-Michael Addition
With the assistance of a basic cyanide ion, the branched α,β-unsaturated ketone proceeds through an intramolecular oxa-Michael addition to form the isoflavonone 20 and regenerate the active catalyst 3. The Gibbs free energy profile and the optimized geometries of involved stationary points along the reaction pathway are displayed in Figures 7, 8, respectively.

First, the deprotonation of the phenolic hydroxyl by the cyanide ion forms a complex 18 (via transition state TS17/18) containing phenol anion, HCN, and (PEt3)3Au cation. The formation of the complex 18 is slightly endergonic, in which the nucleophilicity of the oxygen atom of phenol is increased. Then, the complex 18 could convert into the complex 19 via an intramolecular oxa-Michael addition reaction (via transition state TS18/19). Finally, protonation of the enolate anion center with HCN forms the isoflavonone product 20 and regenerates the active catalyst (PEt3)3AuCN 3. The process described above is exergonic by 10.5 kcal/mol (relative to intermediate 17), and the rate-limiting step is the intramolecular nucleophilic addition reaction of complex 18 with a free energy barrier of 15.0 kcal/mol.

In summary, our calculations reveal that a cyanide ion promoted umpolung hydroacylation/intramolecular oxa-Michael addition mechanism, as shown in Scheme 3, is more favorable than the Au(I)/Au(III) redox mechanism proposed previously. The new pathway contains four stages described above. The overall reaction is exergonic by 41.8 kcal/mol. The hydroacylation of phenyl acetylene is the rate-determining step with a free energy barrier of 27.3 kcal/mol. The free energy barrier is in accord with the experimental fact that the studied reaction takes place at 150°C.

Possible Theoretical Explanations of Experimental Findings
To understand the experimental fact that only the isoflavonone product is observed, we also explored the pathway involving the formation of linear enone intermediate 16-lin (related to the formation of flavanone product) via TS15/16-lin (see Scheme 4). The free energy barrier of this transition state is 31.0 kcal/mol, which is higher than TS15/16-bra by 3.7 kcal/mol. These results indicate that the formation of isoflavonone product is dynamically favorable, which is in accord with the experimental observed regioselectivity.

In addition to PEt3, we also investigated the impact of the ligand on the reactivity of this annulation reaction, including PBu3, and PPh3. Key transition states with relative higher free energies (TS7/10, TS12/13, TS15/16-bra), involved in the pathway, were determined to evaluate the ligand effect. The free energy barriers of these different TSs with different phosphines are listed in Table 1 and the related optimized geometries are presented in Figure S4. The calculated results show that the rate-determining step for the studied reaction is the hydroacylation of phenyl acetylene (TS15/16-bra) with free energy barriers of 27.3 and 32.0 kcal/mol, respectively, when the ligands are PEt3, and PPh3, respectively. However, for the PBu3 ligand, both the HCN-assisted 1,2 H-shift (TS12/13) and hydroacylation of phenyl

**TABLE 1 | Comparison of the free energy barriers (kcal/mol) of different TSs with various ligands (PEt3, PBu3, and PPh3).**

| Ligands | PEt3 | PBu3 | PPh3 | Exp. Yield [%] |
|---------|------|------|------|---------------|
| PEt3    | 25.4 | 24.7 | 27.3 | 36            |
| PBu3    | 22.1 | 25.0 | 24.9 | 78            |
| PPh3    | 29.3 | 28.0 | 31.2 | trace         |

**SCHEME 4 | Two possible pathways for the hydroacylation of phenyl acetylene; Activation free energy barriers (ΔG°) are given in kcal/mol.**
acetylene (TS15/16-bra) may be the rate-determining steps, with comparable barriers (25.0 and 24.9 kcal/mol, respectively). The calculated energy barriers are in qualitative accord with the experimental fact that ligands PEt3 and PBu3 provided moderate to good yield (36 and 78%, respectively), while for the ligand PPh3, only trace amounts of products were observed.

The competing benzocondensation (Wöhler, 1832; Zinin, 1839, 1840) pathway of SA via α-cyano carbanion intermediate 13 (see Scheme 5) is also possible. The free energy barrier of the rate-limiting step in this competing pathway is 24.7 kcal/mol (see Figures S5, S6 for more details). However, this process is almost thermally neutral (slightly exergonic by 0.6 kcal/mol, relative to separated reactants), which is incomparable with the pathway of the annulation of SA and phenyl acetylene (described above, exergonic by 41.8 kcal/mol). This result indicates that the generation of the isoflavanone-type product is thermodynamically much more favorable. Our calculations are in accord with the experimental finding that excessive amount of phenyl acetylene (3-fold amounts of SA) are required to improve the yield of the isoflavanone product.

CONCLUSIONS

We have performed DFT calculations assisted by the MD/CD method to explore the detailed mechanism of the AuCN-catalyzed annulation of salicylaldehyde (SA) and phenyl acetylene. Our calculations reveal that a cyanide ion promoted umpolung hydroacylation/intramolecular oxa-Michael addition mechanism is more favorable than the Au(I)/Au(III) redox mechanism proposed previously. The new proposed pathway contains four stages, as shown in Scheme 3: (1) the association of AuCN with three PEt3 molecule generates an active catalyst (PEt3)3AuCN 3; (2) The nucleophilic attack of the cyanide ion on the carbonyl in SA and subsequent HCN-assisted 1,2 H-shift provide the α-cyano carbanion intermediate 13, which is a key intermediate involved in this mechanism; (3) the hydroacylation of phenylacetylene, followed by the elimination of the cyanide ion, leads to a branched α-β unsaturated ketone 17; (4) the intramolecular conjugate addition of the α,β unsaturated ketone forms the isoflavanone product 20 and regenerates the active catalyst 3. The overall reaction is exergonic by 41.8 kcal/mol. The hydroacylation of phenyl acetylene is the rate-determining step and responsible for the regioselectivity with a free energy barrier of 27.3 kcal/mol at T = 423 K and 1.0 atm in toluene. In the umpolung mechanism, the hydroxyl of SA is found to strongly stabilize the cyanide ion involved intermediates and transition states via hydrogen bond interactions, while the Au(I) ion always acts as a counter cation. Our results are in qualitative accord with the experimental findings. The results provide important insight into Au(I)-catalyzed annulation of SA and aryl acetylenes, which may be useful in designing more effective catalysts for synthesis of heterocyclic frameworks.

DATA AVAILABILITY

All datasets generated for this study are included in the manuscript/Supplementary Files.

AUTHOR CONTRIBUTIONS

The work was completed by cooperation of all authors. MY, GW, JZ, and SL were responsible for the study of concept and design of the project. MY performed corresponding calculations. MY, GW, JZ, and SL drafted and revised the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2019.00557/full#supplementary-material
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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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