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External electric field: a new catalytic strategy for the anti-Markovnikov hydrohydrazination of parent hydrazine†

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The anti-Markovnikov hydroamination reaction is considered to be a particular challenge, and one of the reactants, parent hydrazine, is also regarded as a troubling reagent. In this study, we first studied the hydrohydrazination of parent hydrazine via an effective and green catalyst—external electric field (EEF). The calculation results demonstrated that the anti-Markovnikov and Markovnikov pathways are competitive when there was no catalyst. EEF oriented along the negative direction of the X axis (F_x) accelerated the anti-Markovnikov addition reaction. Moreover, it lowered the barrier height of the first step by 16.0 kcal mol⁻¹ (from 27.8 to 11.8 kcal mol⁻¹) when the field strength was 180 ($\times 10^{-4}$) au. Under the same conditions, the Markovnikov reaction pathway was inhibited, which means that EEF achieved the specificity of hydrohydrazination. The solvents are favorable for the first step addition reaction, particularly the synergy between solvents and F_x lowered the barrier heights by 8.3 (C₆H₆) and 10.7 (DMSO) kcal mol⁻¹ for an $F_x = -60$ ($\times 10^{-4}$) au. Besides, the introduction of the electron-withdrawing substituent (trifluoromethyl) is also a good strategy to catalyze hydrohydrazination, while the electron-donating group (methoxy) is unfavorable.

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Introduction

Nitrogen-containing compounds constitute a useful family of natural products,^{1–3} such as agrochemicals, cosmetics, and pharmaceuticals, and are also important intermediates in numerous industrial processes.⁴ The inspection of the known synthesis methods, such as hydroamination reaction, which is the addition of the N–H bond across an unsaturated C–C bond, is a highly desirable atom-efficient strategy starting from readily accessible materials.^{5–14} Therein, the use of hydrazine as a donor of the N–H bond^{7,15–18} is quite appealing to generate potentially useful products.^{19,20} In 2002, Odom *et al.* reported the first example of the intermolecular hydrohydrazination of 1,1-disubstituted hydrazine to alkynes catalyzed by titanium.²¹ Moreover, increasing number of investigation on the hydrohydrazination of hydrazine were carried out using different metal catalysts.^{9–11,21–34} Among these hydrohydrazination reactions, mono- and di-substituted hydrazines are widely

employed. However, the parent hydrazine (H₂NNH₂) is used rarely despite the potential utility of such a reaction. The possible reasons for hampering the usage of parent hydrazine are as follows: first, hydrazine readily forms Werner complexes, which are usually inert.^{35,36} Second, H₂NNH₂ is known as a strong reducing reagent,^{37,38} which can induce the formation of inactive metal (0) particles.³⁹ Lastly, the resulting products can undergo a metal-mediated N–N bond cleavage to form undesired by-products.^{40–42} In 2010, the disclosed palladium-catalyzed cross-coupling of aryl chlorides and tosylates with hydrazine²⁹ is the only example of the transition-metal-catalyzed functionalization of H₂NNH₂. A few years later, gold-catalyzed hydrohydrazination of alkynes with parent hydrazine was reported.^{30–33}

In general, the hydroamination of a terminal unsaturated C–C bond can take place by two possible pathways, namely^{43–49} the Markovnikov (path M) and the anti-Markovnikov (path AM) addition (Scheme 1). The anti-Markovnikov hydroamination is considered to be a particular challenge because it requires the nucleophilic amine to attack the less electrophilic primary carbon and generate the new [M]–C bond at the more sterically encumbered carbon when the metal catalyst was used.⁴³ However, our calculations revealed that the respective barrier heights of the anti-Markovnikov and Markovnikov hydrohydrazination of phenylacetylene and parent hydrazine are 27.8 and 28.1 kcal mol⁻¹, which are competitive when there was no metal complex, respectively (Fig. 1). It probably benefits from the steric effects in the metal-mediated anti-Markovnikov

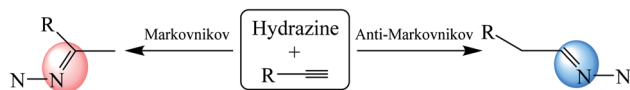
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Scheme 1 Two reaction pathways (Markovnikov: path M and anti-Markovnikov: path AM) for the hydroamination of a terminal alkyne with hydrazine (some of the hydrogen atoms in hydrazine are omitted).

reaction being eliminated. Therefore, the challenging anti-Markovnikov hydrohydrazination of the parent hydrazine becomes a possible pathway. By analyzing the alkyne molecules, we found that reducing the charge of the terminal carbon may be an effective method to generate anti-Markovnikov products because it would make the nucleophilic amine easier to attack the primary carbon atom.

The external electric field (EEF), which is well-known as a green and effective metal-free catalyst, is emerging into one's insight.^{50–53} This “invisible catalyst”, by now, has been investigated by numerous theoretical studies^{54–65} and further confirmed by experiments for the Diels–Alder reaction in 2016.⁶⁶ These investigations manifested that EEF not only influence the atom charge but also catalyzes the chemical reactions. Moreover, using EEF to replace the metal-mediated catalyst could reduce the costs of preparing ligands and metal complexes. Inspired by these reports, we studied the hydrohydrazination of phenylacetylene and parent hydrazine by EEF as a catalyst. The effects of solvents on hydrohydrazination were also taken into account, which are usually considered in experiments. Besides, the local electric field was studied as well, in which the electron-withdrawing and electron-donating groups were used. Another terminal alkyne, namely 1-hexyne, was probed to prove that the obtained conclusion is not occasional. This study conquers the difficulties associated with the usage of parent hydrazine and opens an avenue for extensive applications from theory point of view, and may further provide a new method for the experiments.

Computational details

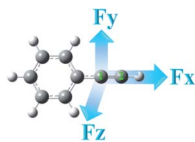
All the studies of the hydrohydrazination of phenylacetylene and parent hydrazine, including geometry optimizations and energy calculations, were carried out using the Gaussian 09 program.⁶⁷ The geometries of the reactants (Rs), transition states (TSs), intermediates (IMs), and products (Ps) were fully optimized at the level of M06/6-31G(d)⁶⁸ because this functional has been proved previously in the hydroamination reactions.^{69–73} All of the stationary points were also calculated by frequency calculations at the same level of theory to confirm that all transition states had only one imaginary frequency, while the others had no imaginary frequency. Furthermore, the transition states were analyzed by intrinsic reaction coordinate (IRC)^{74–76} calculations with the aim of testing the TSs correctly connecting the former and the later stationary points. The single-point energies of all the structures were calculated using M06/6-311++G(d,p). Therefore, the corresponding energies are labelled as M06/6-311++G(d,p)/M06/6-31G(d). Considering the hydrohydrazination reactions are generally performed in solvents, we also examined the solvent effects using the solvation model density (SMD) model⁷⁷ with C₆H₆ ($\epsilon = 2.3$) and DMSO ($\epsilon = 48.9$) solvents. The substituent effects were also considered in this study. The “Field = M \pm N” keyword was used in calculations to probe the effects of external electric fields on hydrohydrazination, which defines the direction and magnitude of EEF in Gaussian 09. In this study, the directions of the used EEF are shown in Scheme 2. The EEF oriented along the C1–C2 triple bond was defined as F_x, with its positive direction pointing from C1 to C2 atom.

Besides, we also employed the M062X method, which combined the other two basis sets (6-311G(d,p) and 6-311++G(d,p)) to prove our calculations. The optimized geometries of every stationary point are similar to those of the previous



Fig. 1 Relative energy (kcal mol⁻¹) profiles for the hydrohydrazination of phenylacetylene and parent hydrazine according to the two pathways.





Scheme 2 Directions (denoted as F_x , F_y , and F_z) of the applied external electric field in the hydrohydrazination of phenylacetylene and parent hydrazine.

method. From the relative energies listed in ESI Tables S2 and S3,[†] we could also obtain the same conclusions as the M06 functional results. Therefore, the calculation results in our study are convincing.

Results and discussions

Anti-Markovnikov hydrohydrazination of phenylacetylene

We first explored the anti-Markovnikov hydrohydrazination of phenylacetylene with parent hydrazine. The optimized geometries of the reactants (R), transition states (TS1 and TS2), intermediate (IM), and product (P) along the potential energy surface with their key bond parameters and relative energies are shown in Fig. 2.

From Fig. 2, it can be seen that when the hydrazine molecule approached the phenylacetylene molecule, a four-membered ring (C1–C2–N4–H3) structure formed, which is the first transition state (TS1) of the anti-Markovnikov hydrohydrazination. The only imaginary frequency of TS1 is $1560.5i \text{ cm}^{-1}$. In TS1, one of the nitrogen atoms (N4) in hydrazine interacted with the terminal carbon atom (C2) of phenylacetylene, accompanied by the formation of a new C2–N4 bond. One of the hydrogen atoms (H3), attached to the nitrogen atom (N4), reacted with the internal carbon atom (C1) in phenylacetylene, and the new C1–H3 bond took its shape. Simultaneously, both the bond lengths of C1–C2 and N4–H3 in TS1 were elongated, implying that the C1–C2 triple bond was impaired, and the N4–H3 bond tended to be ruptured. By inspecting the relative energy of TS1, we could conclude that the barrier height of the first step of this hydrohydrazination is $27.8 \text{ kcal mol}^{-1}$. As the interactions between phenylacetylene and hydrazine were further strengthened, the N4–H3 bond broke completely. Then, the intermediate (IM) of the addition reaction formed, which is also a product of the first addition of N–H to $\text{C}\equiv\text{C}$ bond. The relative energy of IM is

$-37.5 \text{ kcal mol}^{-1}$. Next, the other hydrogen atom (H5) attached to the nitrogen atom (N4) interacted with the internal carbon atom (C1) again, and the second transition state (TS2) of the hydrohydrazination was generated. The sole imaginary frequency of TS2 is $1642.9i \text{ cm}^{-1}$, which connected the intermediate and the product correctly. It can be seen that TS2 is also a four-membered ring structure (C1–C2–N4–H5). Similar to that of TS1, the C1–C2 bond was further impaired, and the distance of N4–H5 was also elongated, tending to break down. Moreover, a new chemical bond, C1–H5, began to form. The interactions between C2 and N4 atoms were further enhanced, along with the corresponding single bond becoming its transitional double bond. From Fig. 2, one can see that the relative energy of TS2 is $12.8 \text{ kcal mol}^{-1}$. Thus, the second addition process needs to overcome a higher barrier of $50.3 \text{ kcal mol}^{-1}$. Ultimately, the N4–H5 bond ruptured, and the final product hydrazone formed.

Hydrohydrazination under external electric field

As mentioned above, the anti-Markovnikov and Markovnikov hydrohydrazination pathways are competitive. Therefore, reducing the barrier heights of the anti-Markovnikov addition is a good strategy to achieve the specific reaction. Based on the fact that the external electric field could change atom charges and catalyze reactions, we first tested the variations of natural bond orbital (NBO) charges on carbon atoms in phenylacetylene under different directions of EEF. The calculated results are shown in Fig. 3. From Fig. 3, it can be seen that EEF oriented along Y and Z axes (Scheme 2) induced little influence on the charges of the C2 atom, while F_x had an obvious effect on it. Therefore, we believe that the EEF oriented along the X-axis is most likely to catalyze the addition reaction. We also calculated the relative energies of anti-Markovnikov hydrohydrazination by the action of all directions of EEF. The results (ESI, Table S5[†]) also verified that F_x could catalyze the reaction much more. On the other hand, it can be seen from Fig. 3 that the charge of the C2 atom decreased with the increase in the external electric field oriented along the negative X-axis. This demonstrated that the nitrogen atoms in hydrazine are more likely to react with the C2 atom, which means the anti-Markovnikov channel is accelerated. On the contrary, the charge of the C2 atom increases gradually when the direction of F_x is flipped, which means that the anti-

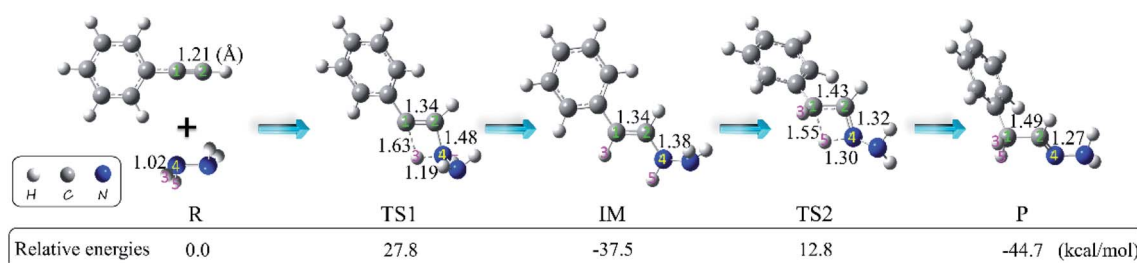


Fig. 2 Geometries (bond lengths are given in Å) and relative energies (in kcal mol^{-1}) of the stationary points in the hydrohydrazination of phenylacetylene with parent hydrazine.



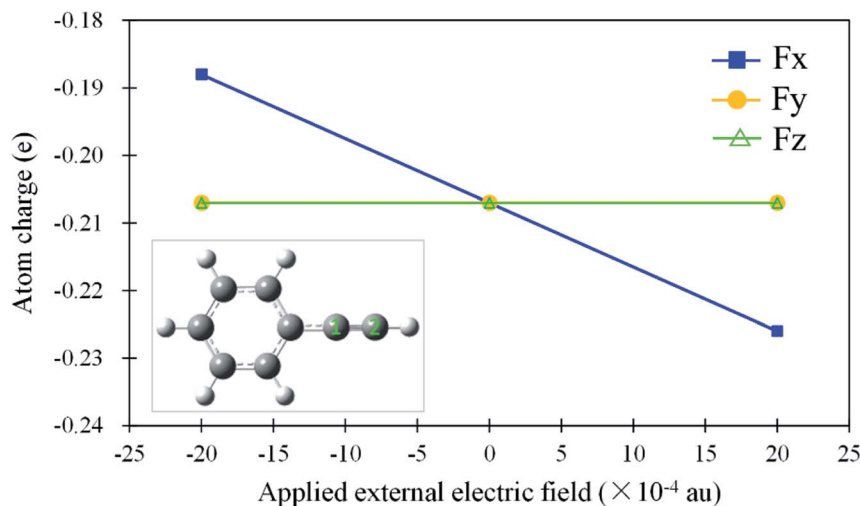


Fig. 3 Variations of the natural bond orbital (NBO) charges (e) on the C2 atom in phenylacetylene under different directions of the external electric field ($\times 10^{-4}$ au).

Markovnikov addition is unfavorable. Consequently, we believe that the negative F_x could accelerate the anti-Markovnikov hydrohydrazination, while the positive F_x inhibits this pathway. In order to prove our assumption, we calculated the effects of both the positive and negative F_x acting on the hydrohydrazination according to path AM and path M. The obtained results in Fig. 4 clearly revealed that the negative F_x decreased the barrier height of path AM and increased the barrier height of path M. In addition, the same direction of F_x has an opposite influence on the two pathways, which also illustrates that it might achieve reaction selectivity. In the following, we focused our study on the effects of the negative F_x for the anti-Markovnikov hydrohydrazination of parent hydrazine.

We re-optimized all the stationary points along the potential energy surfaces at different negative F_x field strengths and recalculated their relative energies. The geometries and energies are shown in Fig. 5 and Table 1, respectively. From Fig. 5, it can be seen that the geometries of the stationary points changed very little under the negative F_x . By inspecting their relative

energies from Table 1, one can conclude that the barrier heights lowered gradually with the increase in field strength. To be specific, for the first step, the barrier height is reduced by $5.7 \text{ kcal mol}^{-1}$ when $F_x = -60 (\times 10^{-4})$ au, equivalent to the rate enhancements by several orders of magnitude. More excitingly, it was lowered by $16.0 \text{ kcal mol}^{-1}$ as the field strength was enhanced to $180 (\times 10^{-4})$ au. For the second step, the influence of negative F_x is also effective but not as large as in the first step. It can be seen that the barrier height of the second addition is lowered by $4.8 \text{ kcal mol}^{-1}$ when $F_x = -180 (\times 10^{-4})$ au. Therefore, the negative F_x had an effective influence on the anti-Markovnikov hydrohydrazination of phenylacetylene with parent hydrazine, particularly for the first step. Why the barrier heights are gradually lowered with the increase in negative F_x , and why the second step had a smaller variation than the first step? We analyzed the frontier molecular orbitals of 'R' (phenylacetylene) and 'IM' (intermediate) under different negative F_x , in which 'R' and 'IM' were separated from their corresponding optimized transition states. In addition, the absolute energies of the two transition states under negative F_x were also

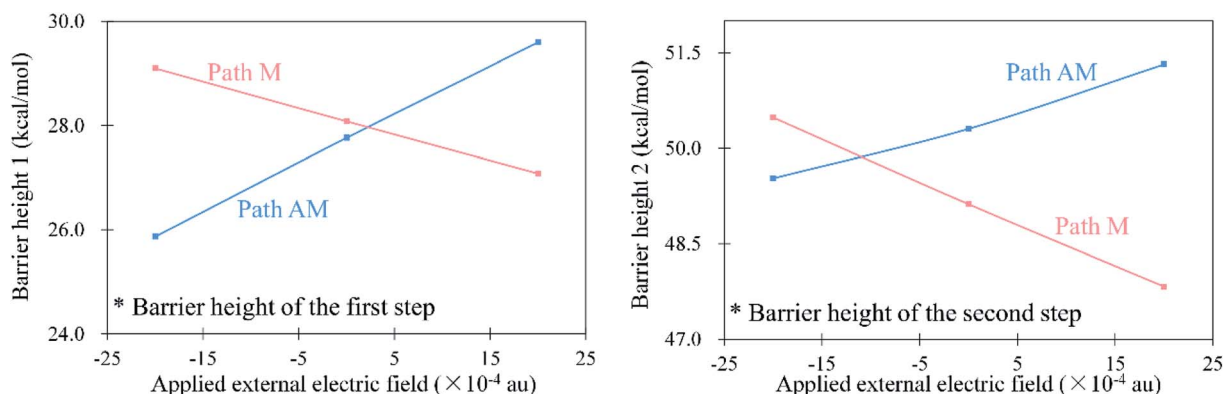

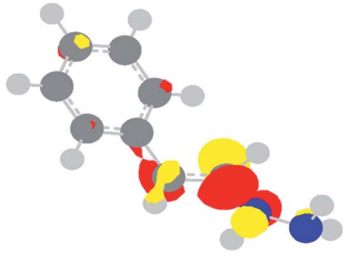
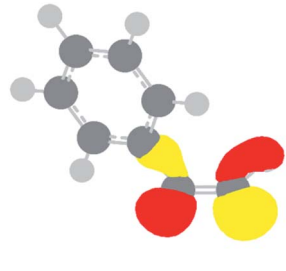




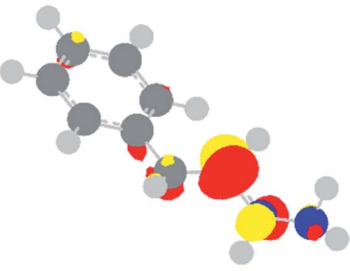


Fig. 4 Barrier heights (kcal mol^{-1}) of the hydrohydrazination according to path AM and path M under the positive and negative F_x ($\times 10^{-4}$ au).



Table 2 Images of the lowest unoccupied molecular orbital (LUMO, isovalue = 0.08) of 'R' (phenylacetylene) and 'IM' (intermediate) and their energies (eV) and the absolute energies (in au) of the transition states (TS1 and TS2) under different negative F_x ($\times 10^{-4}$ au)

| F_x , ($\times 10^{-4}$ au) | LUMO-R, $E_{\text{LUMO-R}}$ (eV) | E_{TS1} (au) | LUMO-IM, $E_{\text{LUMO-IM}}$ (eV) | E_{TS2} (au) |
|--------------------------------|--|-----------------------|---|-----------------------|
| 0 |  -0.0886 | -419.8401 |  -0.0328 | -419.8639 |
| -20 |  -0.0895 | -419.8429 |  -0.0316 | -419.8663 |
| -40 |  -0.0901 | -419.8464 |  -0.0302 | -419.8698 |
| -60 |  -0.0906 | -419.8505 |  -0.0289 | -419.8740 |

further accelerated the addition reactions in the presence of an electron-withdrawing substituent. Besides, the combination of trifluoromethyl phenylacetylene and negative F_x lowered the barrier heights much more than that of phenylacetylene, particularly when the field strength is not large. Therefore, applying the substituted phenylacetylene combined with

a small EEF is a promising method for catalyzing anti-Markovnikov hydrohydrazination.

Hydrohydrazination of 1-hexyne

In order to test that the above-mentioned conclusions are not occasional, the hydrohydrazination of aliphatic 1-hexyne with parent hydrazine was also investigated. The C_6H_6



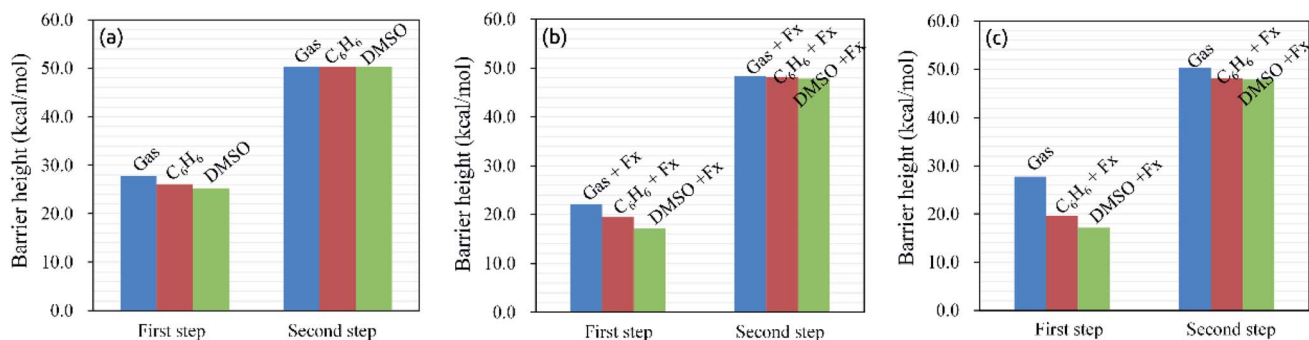


Fig. 6 Barrier heights (kcal mol^{-1}) of hydrohydrazination for the first and second steps in the gas phase, C_6H_6 and DMSO solvents, and negative F_x ($\times 10^{-4}$ au). (a) In solvents; (b) in solvents and negative F_x ; (c) in the gas phase (the blue column) and in solvents and negative F_x (red and green columns).

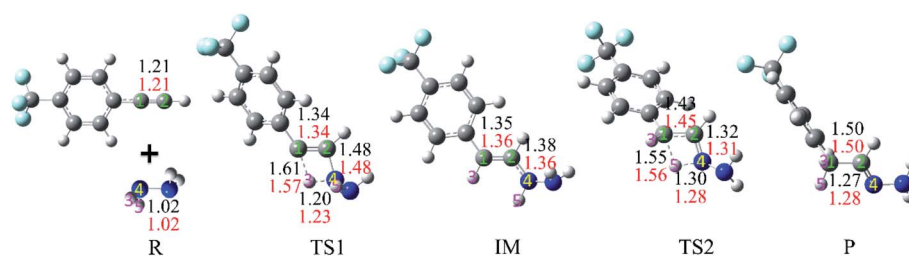


Fig. 7 Geometries (bond lengths are given in Å) of the stationary points in the hydrohydrazination of trifluoromethyl phenylacetylene with parent hydrazine (values in red were calculated with an external electric field, $F_x = -60 \times 10^{-4}$ au).

Table 3 Relative energies (in kcal mol^{-1}) of the stationary points in the hydrohydrazination of the substituted phenylacetylene with parent hydrazine

| | Trifluoromethyl | Methoxy |
|-----|-----------------|---------|
| R | 0.0 | 0.0 |
| TS1 | 25.7 | 30.7 |
| IM | -39.2 | -36.9 |
| TS2 | 10.9 | 13.7 |
| P | -45.4 | -41.6 |

Fig. 9 and 10, respectively. It can be seen that (i) the reaction mode of the hydrohydrazination of 1-hexyne is similar to that of phenylacetylene; (ii) the application of EEF lowered the barrier height of the first step of the reaction; (iii) whether the solvent effect or the introduction of an electron-withdrawing substituent accelerated the hydrohydrazination; (iv) the combination of solvent or substituent and EEF further decreased the reaction barrier heights. All the phenomena are consistent with the hydrohydrazination of phenylacetylene with parent hydrazine.

solvent as well as the substituted alkyne (6-trifluoromethyl-1-hexyne) were considered simultaneously. The obtained geometries and barrier heights of the first step are shown in

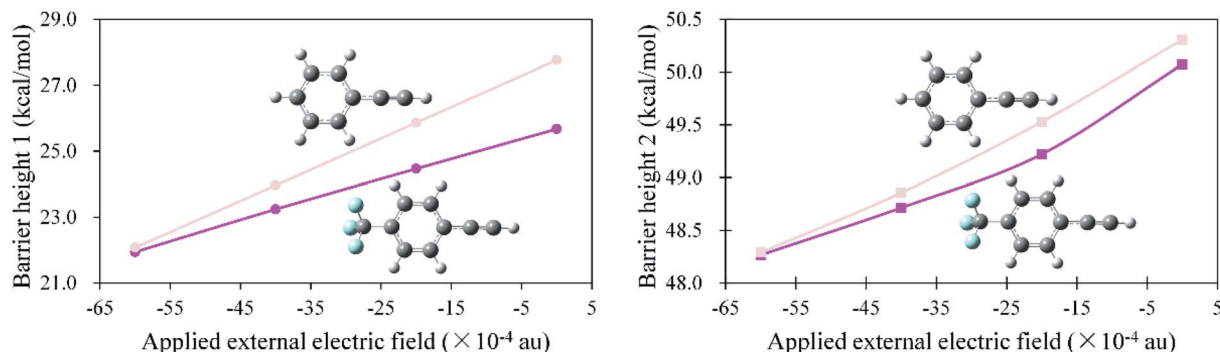


Fig. 8 Barrier heights (kcal mol^{-1}) for the hydrohydrazination of phenylacetylene and trifluoromethyl phenylacetylene with parent hydrazine under negative F_x ($\times 10^{-4}$ au).



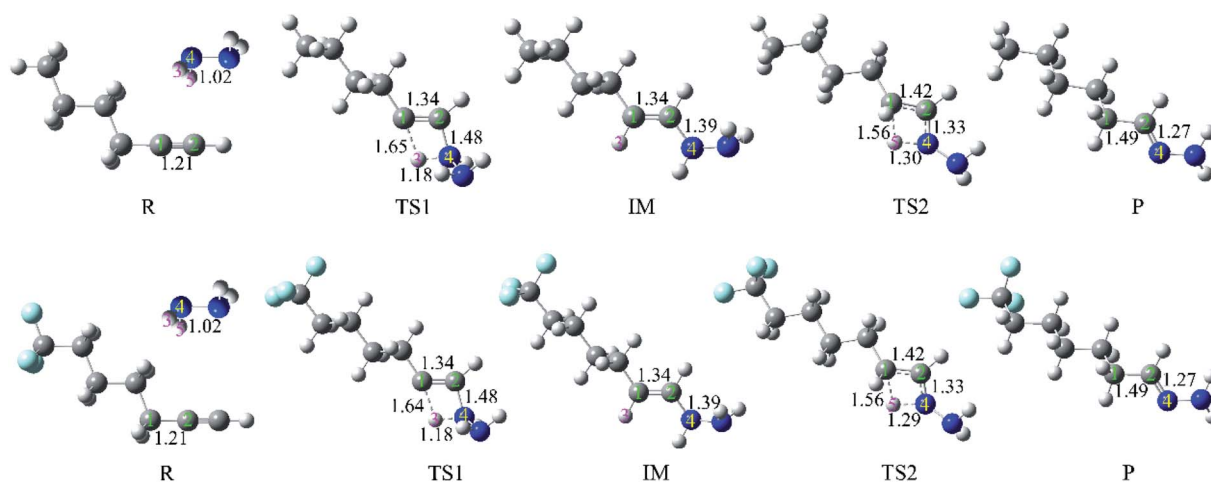


Fig. 9 Geometries (bond lengths are given in Å) of the stationary points in hydrohydrazination of 1-hexyne (6-trifluoromethyl-1-hexyne) with parent hydrazine.

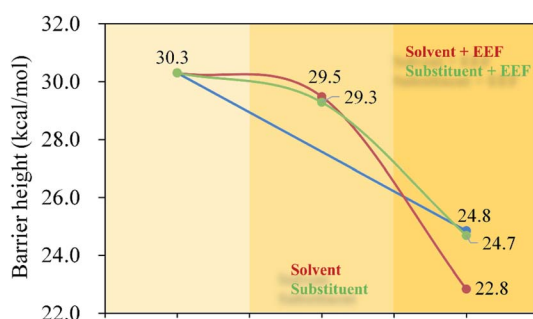


Fig. 10 Barrier heights (kcal mol^{-1}) of the hydrohydrazination of 1-hexyne for the first step in the gas phase, C_6H_6 solvent, and external electric field ($F_x = -60 \times 10^{-4}$ au) and the substituted 1-hexyne.

Conclusions

The effective and green external electric field (EEF) catalyzed anti-Markovnikov hydrohydrazination of parent hydrazine was first studied in the present study using the M06 functional. There are two possible pathways (path M and path AM) in the hydrohydrazination reactions, which are competitive with each other when there was no metal catalyst. Once EEF was introduced, the anti-Markovnikov addition pathway was accelerated when its direction oriented along the negative F_x , which would achieve the selectivity of hydrohydrazination of the parent hydrazine. Furthermore, the larger the field strength of the negative F_x , the lower the barrier heights. The solvents reduced the barrier heights of the first step, but had little effects on the second step. The introduction of substituents in phenylacetylene demonstrated that the electron-withdrawing substituent (trifluoromethyl) could accelerate the reactions. In summary, the best strategy for accelerating this hydrohydrazination is by combining solvent or substituent and external electric field. Hopefully, this study would provide a new guidance for the extensive applications of hydrohydrazination of parent hydrazine in experiments, particularly for the first step of addition reactions.

Conflicts of interest

There are no conflicts to declare.

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