

RESEARCH OUTPUTS / RÉSULTATS DE RECHERCHE

Concerted versus ionic mechanisms of the α and γ extensions in uncatalyzed Mukaiyama reaction between β,γ -unsaturated bis silyl ketene acetal and benzaldehyde

Hadj Mohamed, Slim; Champagne, Benoît

Published in:
Computational and Theoretical Chemistry

DOI:
[10.1016/j.comptc.2021.113395](https://doi.org/10.1016/j.comptc.2021.113395)

Publication date:
2021

Document Version
Publisher's PDF, also known as Version of record

[Link to publication](#)

Citation for pulished version (HARVARD):

Hadj Mohamed, S & Champagne, B 2021, 'Concerted versus ionic mechanisms of the α and γ extensions in uncatalyzed Mukaiyama reaction between β,γ -unsaturated bis silyl ketene acetal and benzaldehyde: A DFT study', *Computational and Theoretical Chemistry*, vol. 1204, 113395.
<https://doi.org/10.1016/j.comptc.2021.113395>

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal ?

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.



Concerted versus ionic mechanisms of the α and γ extensions in uncatalyzed Mukaiyama reaction between β,γ -unsaturated bis silyl ketene acetal and benzaldehyde: A DFT study

Slim Hadj Mohamed^{*}, Benoît Champagne

Laboratory of Theoretical Chemistry, Theoretical and Structural Physical Chemistry Unit, University of Namur, Rue de Bruxelles, 61, B-5000 Namur, Belgium

ARTICLE INFO

Keywords:

DFT
 α adduct Mukaiyama reaction
 γ adduct Mukaiyama reaction
Reaction mechanism
Regioselectivity

ABSTRACT

The uncatalyzed regioselective Mukaiyama aldol reaction between β,γ -unsaturated bis silyl ketene acetal and benzaldehyde has been studied theoretically using density functional theory with the M06-2X exchange–correlation functional. These DFT calculations mostly demonstrate that (i) the α and γ adducts in uncatalyzed Mukaiyama reaction can proceed through two mechanisms, ionic and concerted, (ii) the concerted mechanism is favored for the α adduct and it is disfavored for the γ adduct, (iii) under 12 and 17 kbar, the α adduct reaction path is below the γ adduct reaction path whereas if the pressure decreases to 5 kbar, the opposite is obtained.

1. Introduction

Vinyllogous Mukaiyama aldol reaction [1–10] is the vinyllogous extension of the aldol reaction leading to the formation of C–C bond between β,γ -unsaturated silyl enol ether [11] or silyl ketene acetal [12] and carbonyl compounds to produce linear aldol products through γ extension (Fig. 1). This condensation type was discovered by Mukaiyama and co-workers [1] in 1975, when the reaction was realized between β,γ -unsaturated trialkylsilyl enol-ethers and carbonyl compounds catalyzed by Lewis acid (Fig. 1).

But in the absence of catalyst [2], both linear and branched aldol products will be possible. Indeed, in 1997, Bellassoued et al. [2] studied the uncatalyzed condensation between β,γ -unsaturated bis silyl ketene acetal **1** and benzaldehyde **2** at 65 °C under high pressure conditions and in dichloromethane solution. This reaction gives two regioisomers, one product resulting from a vinyllogous Mukaiyama aldol reaction yielding an γ adduct (linear aldol **3**, Fig. 2) and the second one resulting in an α adduct (branched aldol **4**, Fig. 2). The regioselectivity of this reaction is dependent on pressure. Indeed, the γ -adduct **3** is the major under 2 and 5 kbar but it is the minor under 12 and 17 kbar.

In this manuscript, density functional theory is employed to study the different reaction mechanisms that can take place between **1** and **2** in the absence of a catalyst. This work complements our previous DFT investigation [13] on the uncatalyzed Mukaiyama reaction between C, O,O-tris(trimethylsilyl)ketene acetal and aldehyde, which we

demonstrated that the reaction is possible via two mechanisms, concerted or stepwise, and that the concerted mechanism is favored. The study of the concerted mechanism highlights four types of cyclic transition states (two pro-anti and two pro-syn) [13].

2. Computational methods

Equilibrium structures of reactants and products were optimized at the DFT level using the M06-2X exchange–correlation functional [14,15] and the 6-311G* basis set. The transition states were localized and characterized using the same M06-2X/6-311G* level of approximation. Intrinsic Reaction Coordinate (IRC) calculations were then performed to check that the transition states are related to the corresponding reactants and products. For all species, reactants, products, and transition states, the Gibbs free energy, were evaluated ($T = 338.15$ K, $P = 2, 5, 12$, and 17 kbar) for the reaction between β,γ -unsaturated bis silyl ketene acetal **1** and benzaldehyde **2**. Solvent effects (dichloromethane) were taken into account both in the geometry optimizations and in the calculations of the transition states by using the Integral Equation Formalism (IEF) version of the Polarizable Continuum Model (IEF-PCM) [16,17]. Local nucleophilicity was evaluated using the Fukui function [18]. Calculations were performed using the Gaussian 09 package [19].

^{*} Corresponding author.

E-mail address: slimhadjmohamed4@gmail.com (S. Hadj Mohamed).

<https://doi.org/10.1016/j.comptc.2021.113395>

Received 18 May 2021; Received in revised form 27 July 2021; Accepted 29 July 2021

Available online 8 August 2021

2210-271X/© 2021 Elsevier B.V. All rights reserved.

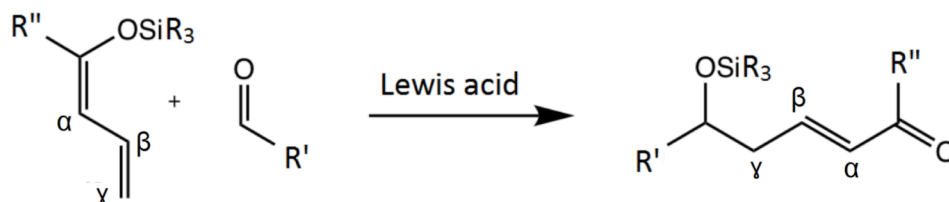
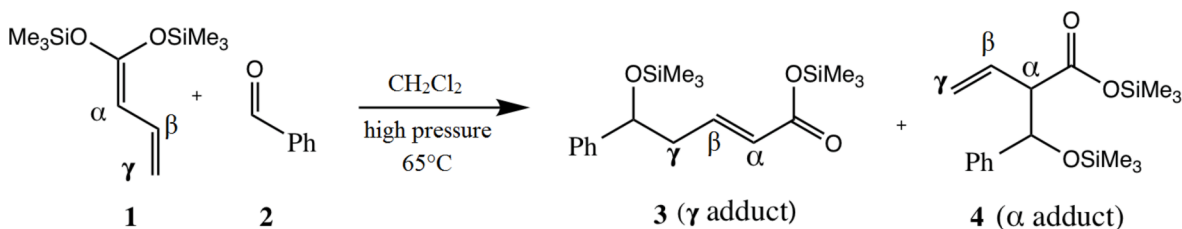


Fig. 1. General scheme for LA-catalyzed Mukaiyama aldol reaction between a β,γ -unsaturated silyl enol ether and an aldehyde.



Entry	Conditions	3	4	Yield %
1	17 kbar, 65°C	25	75	68
2	12 kbar, 65°C	35	65	51
3	5 kbar, 65°C	83	17	57
4	2 kbar, 65°C	88	12	41

Fig. 2. Uncatalyzed Mukaiyama aldol reaction between β,γ -unsaturated bis silyl ketene acetal **1** and benzaldehyde **2**.

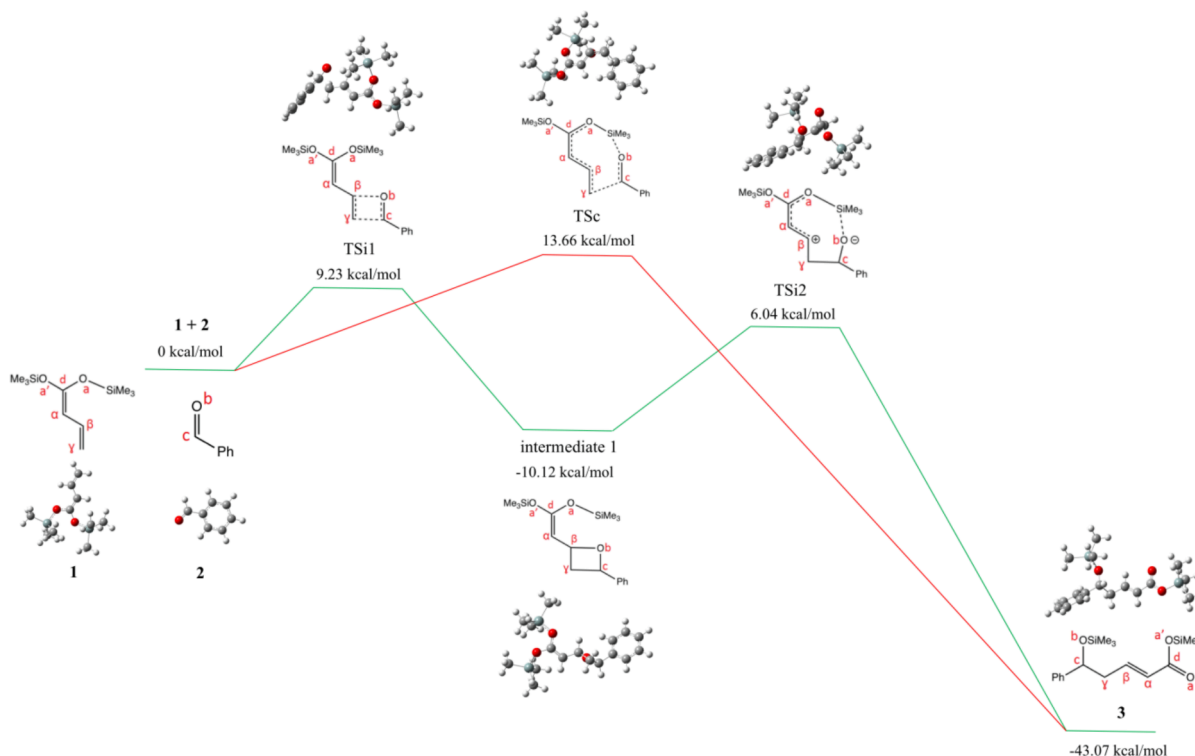


Fig. 3. Energy profile [IEFPCM(dichloromethane)/M06-2X/6-311G(d)] for the ionic versus concerted mechanisms of the γ extension Mukaiyama reaction between β,γ -unsaturated bis silyl ketene acetal **1** and benzaldehyde **2** under 2 kbar.

3. Results and discussion

3.1. Reaction mechanism

The DFT study of the reaction mechanism of the uncatalyzed

vinyllogous Mukaiyama aldol reaction between β,γ -unsaturated bis silyl ketene acetal **1** and benzaldehyde **2** leading to the formation of the regioisomer **3** shows that two reaction paths are possible, concerted and ionic. The concerted mechanism involves the migration of a SiMe₃ group from the O_a oxygen of the acetal to the O_b oxygen of aldehyde together

Table 1

Activation energy (ΔE^\ddagger , kcal/mol), activation enthalpy (ΔH^\ddagger , kcal/mol), and activation free enthalpy (ΔG^\ddagger , kcal/mol) as evaluated with the IEFPCM/M06-2X/6-311G* method (P = 2 kbar, T = 338.15 K, solvent = dichloromethane) for the reaction between β,γ -unsaturated bis silyl ketene and benzaldehyde.

Reaction	TS	ΔE^\ddagger	ΔH^\ddagger	ΔG^\ddagger
γ adduct	TSc	13.66	14.83	30.18
	TSi1	9.23	11.14	27.66
	TSi2	6.04	8.42	22.97
α adduct	TSc'	15.03	16.42	32.31
	TSi1'	18.73	18.93	36.20
	TSi2'	13.31	15.02	34.11

with the formation of the C_c-C_γ bond. The ionic mechanism is characterized by two transition states (TSi1 and TSi2). The first one (TSi1) corresponds to the formation of C_γ-C_c and C_β-O_b single bonds in a concerted process leading to the 4-ring oxetane intermediate (intermediate 1) (Fig. 3). The second (TSi2) leads to the C—O cleavage and migration of the SiMe₃ group from the acetal to the O_b oxygen of benzaldehyde (Fig. 3). Note that no stable [4 + 2] cycloaddition between S-cis conformation of **1** and the C=O carbonyl of **2** has been obtained due to the steric hindrance caused by the geminal OSiMe₃ groups (O_a-SiMe₃ and O_{a'}-SiMe₃) of **1** [20,21]. As shown in Fig. 3 and Table 1, TSi1 lies at lower energy ($\Delta E^\ddagger = 9.23$ kcal/mol, $\Delta H^\ddagger = 11.14$ kcal/mol, $\Delta G^\ddagger = 27.66$ kcal/mol) than the TSc transition state of the concerted mechanism ($\Delta E^\ddagger = 13.66$ kcal/mol, $\Delta H^\ddagger = 14.83$ kcal/mol, $\Delta G^\ddagger = 30.18$ kcal/mol). On the other hand, the energy difference between TSi2 and intermediate 1 is less than 27.74 kcal/mol [E(TSi2) – E(intermediate 1) = 16.16 kcal/mol]. Therefore, the ionic mechanism is favored over the concerted one. These results are the opposite than those of the uncatalyzed α adduct reaction mechanism between **1** and **2** leading to the formation of regioisomer **4**, of which the concerted mechanism is more favorable than the ionic one. Indeed, the following ordering of the transition state energies (E, H, and G) is observed: Δ^\ddagger (TSi1') > Δ^\ddagger (TSc') > Δ^\ddagger (TSi2') (Table 1 and Fig. 4). During the

concerted mechanism, the migration of the silyl group from the silyl ketene acetal **1** to the benzaldehyde **2** is accompanied by the formation of a CC single bond in a boatlike six-membered ring transition state geometry whereas the ionic mechanism is characterized by two transition states (TSi1' and TSi2'), the first one (TSi1') corresponds to the formation of C_α-C_c and C_d-O_b single bonds in a concerted process leading to the intermediate 1' (Fig. 4). The second (TSi2') leads to the migration of the SiMe₃ group to the carbonyl of benzaldehyde to produce the corresponding product (Fig. 4). On the other hand, the same ordering of the transition state energies was obtained when one of the OSiMe₃ group from β,γ -unsaturated bis silyl ketene acetal **1** has been removed to obtain β,γ -unsaturated silyl enol ether [Δ^\ddagger (TSc) > Δ^\ddagger (TSi1) > Δ^\ddagger (TSi2) and Δ^\ddagger (TSi1') > Δ^\ddagger (TSc') > Δ^\ddagger (TSi2')] (Table S1 of supporting information).

3.2. Regioselectivity

The regioselectivity of the uncatalyzed reaction between **1** and **2** is dependent on pressure (Fig. 2). Indeed, the γ adduct is favored under 2 and 5 kbar but it is disfavored under 12 kbar and 17 kbar (Fig. 2). This pressure dependence of the regioselectivity is in agreement with the

Table 2

Local nucleophilicity of C_γ and C_α [N(C_γ)/N(C_α), eV], activation free enthalpy (ΔG^\ddagger , kcal/mol), and activation free enthalpy differences ($\Delta\Delta G^\ddagger$, kcal/mol) between TSi1 and TSc' transition states as evaluated with the IEFPCM/M06-2X/6-311G* method (P = 17, 12, 5, and 2 kbar, T = 338.15 K, solvent = dichloromethane).

Conditions	3/4 ^a	ΔG^\ddagger (TSi1)/ ΔG^\ddagger (TSc')	$\Delta\Delta G^\ddagger$ ^b	N(C _γ)/N(C _α)
17 kbar, 338.15 K	25/75	30.85/28.93	1.92	0.89/0.96
12 kbar, 338.15 K	35/65	30.76/29.51	1.25	0.72/0.76
5 kbar, 338.15 K	83/17	27.70/32.22	– 4.52	0.49/0.38
2 kbar, 338.15 K	88/12	27.66/32.31	– 4.65	0.43/0.30

^b $\Delta\Delta G^\ddagger = \Delta G^\ddagger$ (TSi1) – ΔG^\ddagger (TSc').

^a 3/4 ratio obtained by Bellassoued et al. [2].

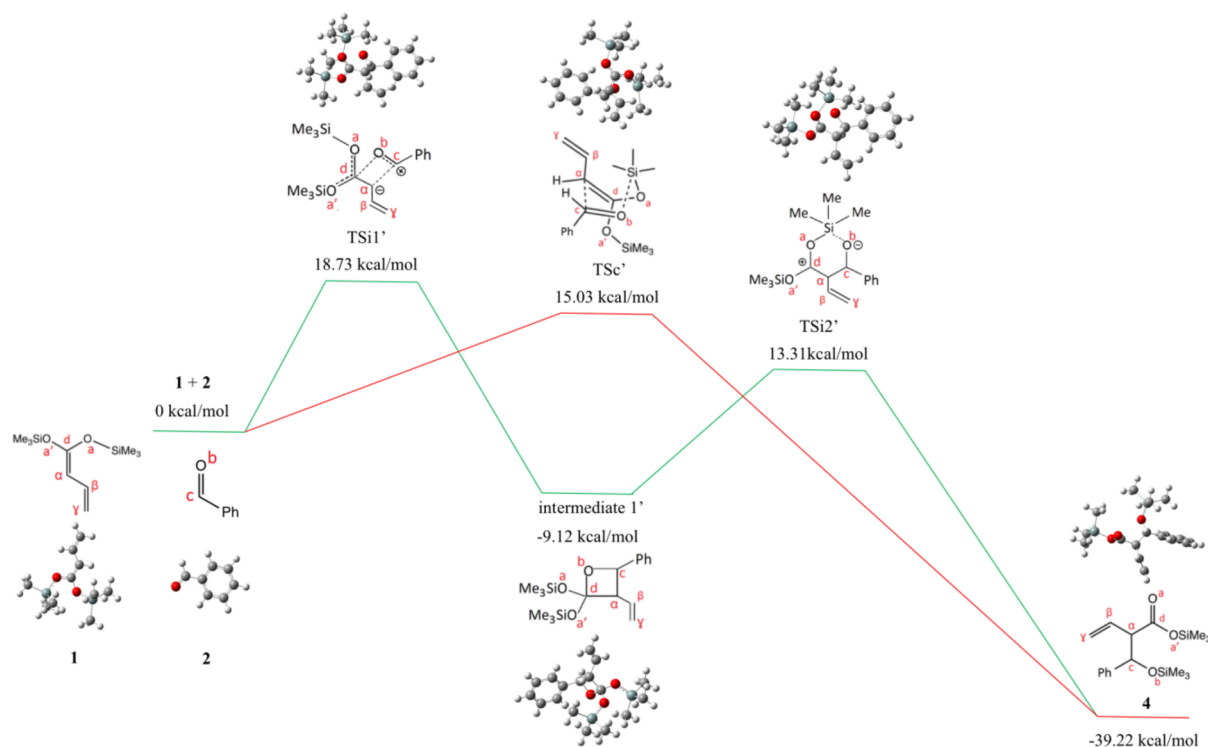


Fig. 4. Energy profile [IEFPCM(dichloromethane)/M06-2X/6-311G(d)] for the ionic versus concerted mechanisms of the α extension Mukaiyama reaction between β,γ -unsaturated bis silyl ketene acetal **1** and benzaldehyde **2** under 2 kbar.

activation free enthalpy differences between TS1 and TS γ transition states [ΔG^\ddagger (TS1) < ΔG^\ddagger (TS γ) under 2 and 5 kbar; ΔG^\ddagger (TS1) > ΔG^\ddagger (TS γ) under 12 and 17 kbar] (table 2) and the local nucleophilicity values for α and γ carbons of β,γ -unsaturated bis silyl ketene acetal [N(C γ) > N(C α) under 2 and 5 kbar; N(C γ) < N(C α) under 12 and 17 kbar] (table 2).

4. Conclusion

In 1997, Bellassoued and co-workers [2], studied the Mukaiyama aldol reaction between β,γ -unsaturated bis silyl ketene acetal **1** and benzaldehyde **2** under high pressure conditions to produce γ -adduct **3** and α -adduct **4** regioisomers. Our DFT calculation with the M06-2X exchange–correlation functional shows that (i) this reaction takes place via a cyclic transition states, (ii) the ionic mechanism is favored for the γ -adduct reaction (**1** + **2** \rightarrow TS1 \rightarrow intermediate **1** \rightarrow TS2 \rightarrow **3**) while the concerted mechanism is favored for the α -adduct reaction (**1** + **2** \rightarrow TS γ \rightarrow **4**), (iv) The regioselectivity of the reaction can be explained by comparing the TS1 and TS γ free energies and the local nucleophilicity values of C γ and C α atoms.

CRedit authorship contribution statement

Slim Hadj Mohamed: Conceptualization, Methodology, Software, Data curation, Writing - original draft, Visualization, Investigation, Software, Writing - review & editing. **Benoît Champagne:** Supervision, Validation.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.comptc.2021.113395>.

References

- [1] T. Mukaiyama, A. Ishida, A new synthesis of vitamin A, *Chem. Lett.* 4 (11) (1975) 1201–1202.
- [2] M. Bellassoued, E. Reboul, F. Dumas, High Pressure Induced Mukaiyama Type Aldol Reaction of bis Trimethylsilyl Ketene Acetals, *Tetrahedron Lett.* 38 (32) (1997) 5631–5634.

- [3] K. Furuta, T. Maruyama, H. Yamamoto, Catalytic asymmetric aldol reactions. Use of a chiral (acyloxy)borane complex as a versatile Lewis-acid catalyst, *J. Am. Chem. Soc.* 113 (3) (1991) 1041–1042.
- [4] C. Kaneko, M. Sato, S. Sunami, Y. Sugita, An efficient asymmetric aldol reaction of 4-trimethylsiloxy-6-methylene-1,3-dioxines by chiral binaphthol-titanium complex catalysis, *Heterocycles* 41 (7) (1995) 1435, <https://doi.org/10.3987/COM-95-7065>.
- [5] M. Christmann, M. Kalesse, Vinylogous Mukaiyama aldol reactions with triarylboranes, *Tetrahedron Lett.* 42 (7) (2001) 1269–1271.
- [6] J. Hassfeld, M. Christmann, M. Kalesse, Rapid Access to Polyketide Scaffolds via Vinylogous Mukaiyama Aldol Reactions, *Org. Lett.* 3 (22) (2001) 3561–3564.
- [7] J. Hassfeld, M. Kalesse, Advances in the vinylogous Mukaiyama aldol reaction and its application to the synthesis of the C1–C7 subunit of oleandolide, *Tetrahedron Lett.* 43 (29) (2002) 5093–5095.
- [8] R. Villano, M.R. Acocella, M. De Rosa, A. Soriente, A. Scettri, Pronounced asymmetric amplification in the aldol condensation of Chan's diene promoted by a Ti (IV)/BINOL complex, *Tetrahedron Asymmetry* 15 (2004) 2421–2424.
- [9] C.S. López, R. Álvarez, B. Vaz, O.N. Faza, Á.R. de Lera, Simple Diastereoselectivity of the BF $_3$ ·OEt $_2$ -Catalyzed Vinylogous Mukaiyama Aldol Reaction of 2-(Trimethylsiloxy)furan with Aldehydes, *J. Org. Chem.* 70 (9) (2005) 3654–3659.
- [10] C. Curti, L. Battistini, F. Zanardi, G. Rassu, V. Zambrano, L. Pinna, G. Casiraghi, Uncatalyzed Diastereoselective Vinylogous Mukaiyama Aldol Reactions on Aqueous Media : Pyrrole vs Furan 2-Silyloxy Dienes, *J. Org. Chem.* 75 (2010) 8681–8684.
- [11] H.O. House, L.J. Czuba, M. Gall, H.D. Olmstead, The Chemistry of carbanions. XVIII. Preparation of Trimethylsilyl Enol Ethers, *J. Org. Chem.* 34 (8) (1969) 2324–2336.
- [12] C. Ainsworth, Y. Kuo, Ketene bis(trialkylsilyl) acetals: synthesis, pyrolysis and spectral studies, *J. Organometal. Chem.* 46 (1972) 73–87.
- [13] S. Hadj Mohamed, M. Trabelsi, B. Champagne, Unraveling the Concerted Reaction Mechanism of the Noncatalyzed Mukaiyama Reaction between C, O, O-Tris(trimethylsilyl)ketene Acetal and Aldehydes Using Density Functional Theory, *J. Phys. Chem. A* 120 (28) (2016) 5649–5657.
- [14] Y. Zhao, D.G. Truhlar, Comparative DFT study of van der Waals complexes : rare-gas dimers, alkaline-earth dimers, zinc dimer, and zinc-rare-gas dimers, *J. Phys. Chem. A* 110 (2006) 5121–5129.
- [15] Y. Zhao, D.G. Truhlar, The M06 suite of density functionals of main group thermochemistry, thermochemical kinetics, noncovalent interactions, excited states, and transition elements: two new functional and systematic testing of four M06-class functionals and 12 other functionals, *Theor. Chem. Acc.* 120 (2008) 215–241.
- [16] D.M. York, M. Karplus, A Smooth Solvation Potential Based on the Conductor-Like Screening Model, *J. Phys. Chem. A* 103 (50) (1999) 11060–11079.
- [17] J. Tomasi, B. Mennucci, R. Cammi, Quantum mechanical continuum solvation models, *Chem. Rev.* 105 (8) (2005) 2999–3094.
- [18] S. Deuri, P. Phukan, A density functional theory study on π -nucleophilicity and electron-transfer oxidation of silyl enol ethers and ketene silyl acetals, *J. Mol. Struct. (Theochem)* 945 (1–3) (2010) 64–70.
- [19] M.J. Frisch, G.W. Trucks, H.B. Schlegel, G.E. Scuseria, M.A. Robb, J.R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G.A. Petersson, H. Nakatsuji, et al. Gaussian 09, revision A.01; Gaussian, Inc.: Wallingford, CT, (2010).
- [20] S.H. Mohamed, B. Champagne, M. Trabelsi, DFT Investigation of the Diastereoselectivity of the MX $_2$ and MX $_3$ Lewis-Acid-Catalyzed Mukaiyama Aldol Reaction between C, O, O-Tris(trimethylsilyl)ketene Acetal and Aldehydes, *J. Phys. Chem. A* 122 (8) (2018) 1938–1947.
- [21] J.M. Lee, P. Helquist, O. Wiest, Diastereoselectivity in Lewis-Acid-Catalyzed Mukaiyama Aldol Reaction: A DFT Study, *J. Am. Chem. Soc.* 134 (2012) 14973–14981.