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Revisiting oxime–nitron tautomerism. Evidence of nitron tautomer participation in oxime nucleophilic addition reactions†

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The oxime–nitron tautomerism has been revisited using high-level DFT calculations. The isomerization has been found to be more favorable through a bimolecular process involving two molecules of oxime, a finding that argues against the commonly accepted thermal 1,2-H-shift mechanism. The reaction of arylamidoximes with 1,2-diaza-1,3-dienes to yield the corresponding *O*-substituted oximes (stable intermediates for the synthesis of 1,2,4-oxadiazine derivatives) was also investigated as a rare case in which *O*-alkylation is observed in the reaction between oximes and electron-poor alkenes in the absence of a base. Under such conditions the reaction usually proceeds through the nucleophilic attack of the oxime nitrogen to yield the corresponding nitron. The computational investigation revealed that in the case of arylamidoximes, the pathway involving the less stable but more reactive nitron tautomer is the predominant mechanism, evidencing for the first time the involvement of a nitron tautomer in nucleophilic additions of oximes. Validation of the model has been carried out by studying alternative ene-like processes; the dramatically different reactivity predicted for arylamidoximes and unsubstituted oxime are rationalized in terms of steric hindrance.

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Introduction

About 40 years ago, Vijfhuizen and Terlouw reported the first evidence, observed by mass spectrometry, of the oxime–nitron isomerization (Scheme 1).¹ Two years later, Dignam and Hegarty studied isomerization of oximes in aqueous solution as a function of the pH and concluded that at high pH values the reactive species is the neutral oxime rather than the nitron tautomer.² In 1984, Grigg and co-workers proposed that isomerization of oxime to nitron consisted of a thermal 1,2-hydrogen shift.³

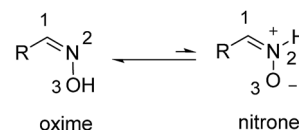
Since these early reports, cycloaddition reactions of oximes towards isoxazolidines have been extensively studied.⁴ In particular, the intramolecular version of the reaction (the so-called “intramolecular oxime olefin cycloaddition”, IOOC) has received considerable attention⁵ in contrast to intermolecular

reactions which have been less studied.⁶ In this context, Tamura and co-workers developed intra-⁷ and intermolecular⁸ cycloadditions of *O*-silylated oximes that required milder conditions by using boron trifluoride etherate as a promoter of the reaction, leading to *N*-boranonitrones. Frank and co-workers reported a combined experimental and computational study demonstrating that even though the nitron tautomer is less stable than the oxime one, the former is more reactive, leading to more stable transition structures.⁹ All these studies concern cycloaddition reactions of the nitron tautomer. However, there are no reports illustrating the participation of such a tautomer in nucleophilic additions in spite of the fact that nitrones can act as excellent electrophiles.¹⁰

The mechanism of the isomerization has been discussed in various studies,¹¹ all of them claiming for the previously proposed 1,2-hydrogen shift. Noguchi and co-workers¹² found a dependence on the type of solvent but no evidences on the effect of alcoholic solvents. The authors suggested a solvation effect favoring the isomerization towards nitron. The influence of substituents in the tautomerism has also been considered¹³

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† Electronic supplementary information (ESI) available: Details on calculations corresponding to oxime–nitron tautomerism including unimolecular 1,2-H shift direct and promoted by neighboring groups; bimolecular mechanisms promoted by EtOH and *via* oxime dimer complexes. Conformational analysis and reactivity indices of 1,2-diaza-1,3-diene **6**. Details on calculations corresponding to Michael addition of oximes **1a** and **1b** to **6**. Details on calculations corresponding to ene-like processes between oxime **1a** and **6**, and oxime **1b** and methyl 2-methylfumarate and **6**. Cartesian coordinates of optimized structures. See DOI: 10.1039/c6ra02638a



Scheme 1 Oxime–nitron tautomerism.

and in some particular case, a facile isomerization has been attributed to the presence of functional groups facilitating hydrogen shift.¹⁴ For instance, for 4-chlorobenzaldehyde it had been estimated that *ca.* 1–2% of the NH-nitrone should be present in equilibrium.¹⁵ Radom and co-workers studied computationally nitrosomethane and its nitron and oxime isomers.¹⁶ These authors concluded that NH-nitrones should be observable even though a barrier of 70.8 kcal mol^{−1} (MP3/6-31G++) was estimated for the 1,2-H shift between formaldoxime and formaldonitrone. A further computational study on the oxime–nitron tautomerism was carried out by Lammertsma and co-workers,¹⁷ who demonstrated that the oxime–nitron tautomerism is thermodynamically unfavoured. More recently, Sousa and co-workers have suggested that the population of the nitron tautomer should be insignificant on the basis of more accurate calculations.¹⁸ However, these recent studies were only concerned with the stability of the different tautomers as well as solvent effects.

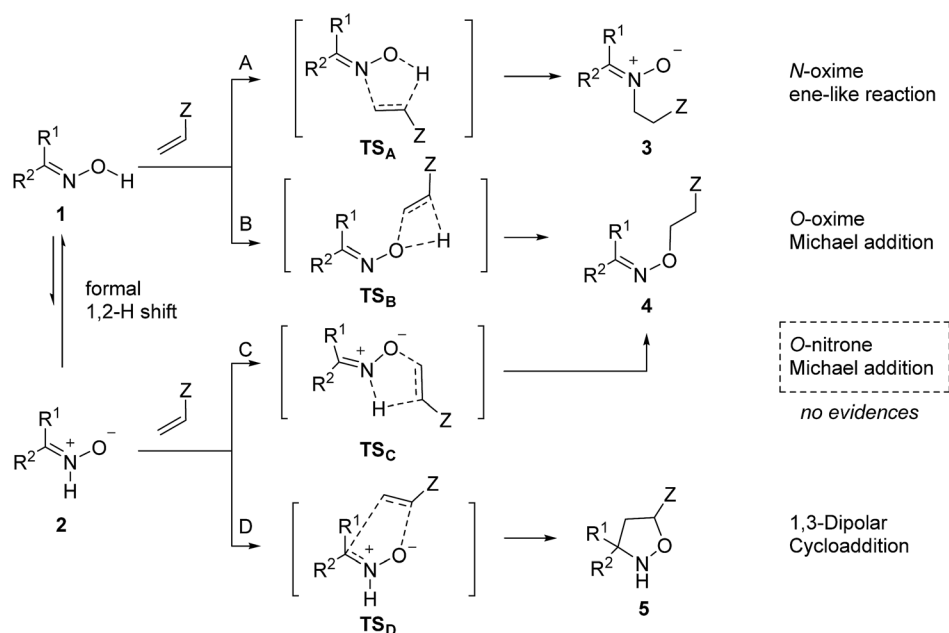
The oxime–nitron tautomerism has important implications in reactivity. When an oxime reacts with an alkene (that can act as both dipolarophile and Michael acceptor if Z is an electron-withdrawing group) three products can be obtained, even though there are four possible reaction pathways (Scheme 2).

When oximes react with electron-poor alkenes (Z = EWG), in the absence of any base, the usually preferred pathway is an ene-like reaction (formal Michael addition) *via* the nitrogen lone pair to yield *N*-alkylnitrones (Scheme 2, path A).^{6a} The mechanism is expected to be concerted (through **TS_A**) with concomitant nucleophilic attack of the nitrogen atom and H-transfer. In fact, the concertedness of the reaction has already been theoretically demonstrated for related additions of hydroxylamines to electron-poor alkenes.¹⁹ Usually, the *in situ* formed nitrones **3** react with a second molecule of alkene to give the

corresponding dipolar cycloaddition (not shown in Scheme 2). On the contrary, when the alkene is unactivated, path D is preferred. As mentioned above, the cycloaddition reaction of the nitron tautomer to give isoxazolidines **5** (Scheme 2, path D), through a typical concerted transition state **TS_D**, has also been reported in several instances.⁴ In the case of intramolecular reactions a high dependence of the stereochemistry of the oxime has been observed; whereas *E*-isomers of oximes react through path A, *Z*-isomers prefer path D *via* tautomeric nitron.^{13c}

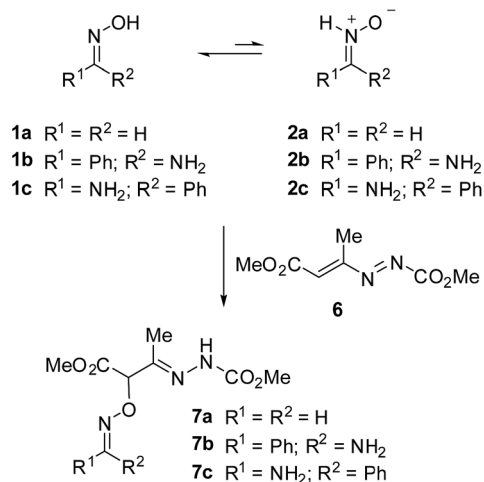
Under basic conditions it is also possible to carry out the *O*-alkylation of the oxime through a typical oxa-Michael reaction.²⁰ However, in the absence of a base the mechanism should take place through a disfavored (due to ring strain) transition state (**TS_B**) leading to *O*-alkylated derivatives **4** (Scheme 2, path B). To date no evidences have been found of a fourth pathway C consisting of the formation of oxime ethers by nucleophilic attack of the nitron oxygen, after isomerization of oxime into nitron. Only two reports described the obtention of *O*-alkylated oximes by reaction with electron-poor alkenes in the absence of a base.^{4i,21} By analogy to path A (**TS_A**) the mechanism could be expected to be concerted through **TS_C**. Although it cannot be considered an ene-like reaction because the hydrogen is not allylic, the process involves a 1,4-H shift as in the case of path A.

As part of our ongoing interest in mechanistic aspects of nitron reactivity,²² we sought to investigate the nitron–oxime isomerization by computational DFT methods and explore the possibility of a participation of the nitron tautomer in nucleophilic addition reactions in the absence of a base. We chose for the study the Michael addition of arylamidoximes to 1,2-diaza-1,3-dienes, studied experimentally by Santeusano and co-workers (Scheme 3).²¹ This reaction, carried out in the absence of a base, is a rare example of addition of oximes to electron-



Scheme 2 Possible reaction pathways between oximes and electron-poor alkenes.





Scheme 3 Nucleophilic addition of oximes to 1,2-diaza-1,3-dienes.

poor olefins in which *O*-alkylated oximes **4** are preferentially formed instead nitrones **3**.

Under these circumstances, paths B and C could be competitive, the tautomerism prior the addition being an option. Therefore, aiming to obtain a more accurate description of the oxime–nitron tautomerism and its implication on nucleophilic addition reactions, this paper presents proofs for a bimolecular isomerization mechanism and, for the first time, an unequivocal characterization of transition state structures in solution involving nitron tautomers in a nucleophilic addition reaction (Scheme 2, path C).

Computational methods

All of the calculations were performed using the Gaussian09 program.²³ Molecular geometries were optimized with the M06-2X functional²⁴ in conjunction with cc-pVTZ basis set.²⁵ This method has been recently used successfully in theoretical investigations with nitrones.^{22a-f} Analytical second derivatives of the energy were calculated to classify the nature of every stationary point, to determine the harmonic vibrational frequencies, and to provide zero-point vibrational energy corrections. The thermal and entropic contributions to the free energies were also obtained from the vibrational frequency calculations, using the unscaled frequencies. All transition structures were characterized by one imaginary frequency. All the located TSs were confirmed to connect to reactants and products by intrinsic reaction coordinate (IRC) analyses.²⁶ The IRC paths were traced using the second order González–Schlegel integration method.²⁷ Calculations have been carried out considering solvent effects (EtOH) with the PCM.²⁸

We have considered two models for the study of oxime–nitron isomerization: (i) the simplest oxime **1a** and nitron **2a**, and (ii) the real system studied experimentally,²² consisting of oxime (*Z*)-**1b** and nitron (*Z*)-**2b**. For the purpose of comparison, we have also included in the study the (*E*)-isomers **1c** and **2c** (Scheme 3).

Results and discussion

Starting materials

We first computed **1a** and both *E/Z* isomers of oximes **1b, c** and nitrones **2b, c**; since the reaction was carried out in EtOH, we chose this solvent for our studies. In the case of oximes, the *anti* conformer was the most stable; in fact, the corresponding *syn* oximes could not be located because all calculations converged to the most stable *anti* oximes. The optimized geometries of compounds **1a–c** and **2a–c** are given in Fig. 1.

Calculations predict the *Z*-isomer to be more stable than the *E*-isomer. In the case of oximes, isomer (*Z*)-**1b** is 5.8 kcal mol^{−1} more stable than (*E*)-**1c**; for nitrones (*Z*)-**2b** is 6.9 kcal mol^{−1} more stable than (*E*)-**2c**. Both (*Z*)-isomers present a hydrogen bond between the amino group and the oxygen atom ($d = 2.15$ Å and 2.21 Å for **1b** and **2b**, respectively). A weaker interaction ($d = 2.45$ Å) is also observed for (*E*)-**1c** between the amino group and the oxime nitrogen. All *Z* and *E* isomers **1b, c** and **2b, c** deviate the phenyl ring from planarity because of the steric hindrance of the phenyl ring. Larger torsion angles are present for (*E*)-isomers than for (*Z*)-isomers ($C_{Ar2}-C_{Ar1}-C_{ox}-N_{ox}$: 53.8° for (*E*)-**1c** vs. 27.5° for (*Z*)-**1b**. $C_{Ar2}-C_{Ar1}-C_{nit}-N_{nit}$: 35.0° for (*E*)-**2c** vs. 26.4° for (*Z*)-**2b**). When compared oxime **1b** and nitron **2b**, the former resulted to be more stable by 9.2 kcal mol^{−1} in agreement with experimental observations. The *E/Z* isomerization of oximes and nitrones has been studied in previous reports and it is well document that it can take place through energy barriers of *ca.* 10 kcal mol^{−1} for oximes²⁹ and 25 kcal mol^{−1} for nitrones.^{21b}

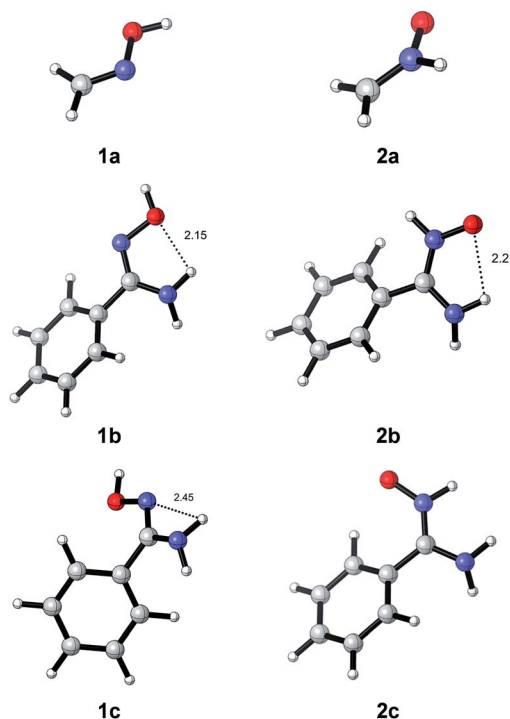


Fig. 1 Optimized transition structures (M06-2X/cc-pVTZ/PCM = EtOH) of oximes **1a–c** and nitrones **2a–c**. Distances are given in angstrom.



Oxime–nitrone tautomerism

Grigg and co-workers initially suggested that 1,2-prototropy in oximes should be a facile process.³⁰ On the other hand, the same authors also mentioned that it is a high energy process when compared with other reactions such as formal Michael additions (actually, ene-like reactions),^{6a} thus explaining the observed behavior in the presence of electron-poor olefins (Scheme 2, path A). With other olefins the isomerization is well demonstrated by trapping the NH-nitrone through a dipolar cycloaddition (Scheme 2, path D).⁴ Since the early proposal of the 1,2-prototropic route by Grigg and co-workers,³ all authors have assumed this rationale¹¹ in spite of such isomerization mechanism had been discarded by Radom and co-workers 30 years ago.¹⁶ In fact, that study considered location of transition structure **TS1a** for the 1,2-H shift between **1a** and **2a** (Scheme 4) and assigned a barrier to the process of 70.8 kcal mol^{−1}.¹⁶ We also located **TS1a** in gas phase (Table 1) and considering ethanol as solvent at our standard level of theory (M06-2X/cc-pVTZ). The results for the reaction in EtOH are summarized in Table 2. The geometry of **TS1a** is given in Fig. 2.

The calculated barrier for the isomerization in gas phase was 54.7 kcal mol^{−1}. The observed difference of *ca.* 20 kcal mol^{−1} with the value obtained by Radom and co-workers in the 80's,¹⁶ can be attributed to the use of M06-2X functional which slightly underestimate activation parameters.³¹ In any case the obtained results give a good illustration of the reaction mechanism pointing out that the process would require a high energy to take place.

When considering solvent effects (EtOH) the barrier for **TS1a** (53.2 kcal mol^{−1}) is similar to that found in gas phase. The corresponding **TS1b** starting from **1b** to form **2b** was also located and a barrier of 49.5 kcal mol^{−1} was found (Table 2). The transition structure (**TS1c**) corresponding to the transformation of less stable **1c** to form **2c** was also located and a higher barrier (56.4 kcal mol^{−1}) was obtained for the process. In any case, high energies (*ca.* 50 kcal mol^{−1}) are required by the process to take place.

The geometries of the transition structures are given in Fig. 2. Similar distances for hydrogen shift are found in all cases. The N–H distances (1.11–1.13 Å) are shorter than the O–H distances (1.28–1.31 Å). The 1,2-H shift takes place in the plane of the oxime for **TS1a** and **TS1c** (H–O1–N2–C3 dihedral angle of 180.0° and 174.7° for **TS1a** and **TS1c**, respectively). On the other hand for **TS1b** the 1,2-H shift takes place out of the plane of the oxime (H–O1–N2–C3 dihedral angle of 132°). This deviation is due to the existing H-bond interaction between an oxygen lone

Table 1 Calculated absolute (hartrees) and relative (kcal mol^{−1}) energies in gas phase of the stationary points corresponding to isomerization of **1a** into **2a** via 1,2-H shift

	M06-2X/cc-pVTZ		MP3/6-31G++ ^a	
	<i>G</i>	ΔG^b	<i>G</i>	ΔG^b
1a	−169.787344	0.0	−169.34926	0.0
TS1a	−169.700184	54.7	−169.23702	70.8
2a	−169.765342	13.8	−169.31409	22.2

^a Data from ref. 16. ^b Referred to **1a**.

pair and the amino group, which is not present in **TS1c**. Whereas **TS1a** is, obviously, completely planar, in **TS1b** and **TS1c** the phenyl ring is out the plane formed by the nitrone and amino moieties (by 25° and 43° in **TS1b** and **TS1c**, respectively).

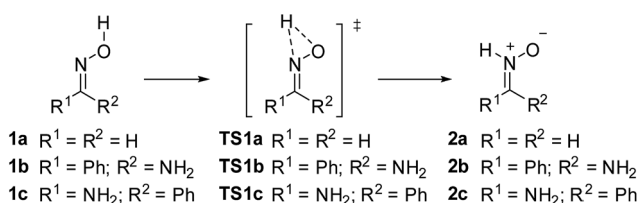
It has been suggested that neighboring groups at the periphery of the oxime could promote a more favorable 1,5-H shift in detriment of the assumed 1,2-H shift. This possibility has been exemplified with monoximes of 1,2,3-triketones,³² alk-2-enylamino-substituted pyrans¹² and pyrido[1,2-*a*]pyrimidines.^{13a,14} In the case of oxime **1b** we also explored a similar option by considering the intramolecular protonation of the amino group and further 1,3-H sigmatropic shift (Scheme 5).

According to the calculated mechanism, illustrated in Scheme 5, the isomerization should take place starting from oxime **1b**. The initial H-transfer from oxime to α -amino group takes place along the reaction path in an early stage without formation of any stationary point. The first transition state found in the reaction path corresponds to rotation of the C–N bond. The IRC showed a temporary pyramidalization of C atom with concomitant transformation of the C=N–O[−] system into a [−]C–N=O one during the rotation (see ESI†). This rotation is required to achieve the correct orientation of the nitrogen lone pair to receive the proton from the amino group in the second step. In fact, such geometry is that found in intermediate **8** whose zwitterionic nature was confirmed by its large dipole moment (11.6 D). The formation of intermediate **8** is the rate-limiting step. Further evolution of **8** takes place through a 1,3-H

Table 2 Calculated (M06-2X/cc-pVTZ/PCM = EtOH) absolute (hartrees) and relative (kcal mol^{−1}) energies of the stationary points corresponding to isomerization of **1a–c** into **2a–c** via 1,2-H shift

	ΔE_0^a	ΔG^a
1a	0.0	0.0
TS1a	53.1	53.2
2a	9.8	10.0
1b	0.0	0.0
TS1b	49.6	49.5
2b	9.1	9.2
1c	6.2	5.8
TS1c	56.8	56.4
2c	15.9	16.1

^a Referred to **1a** or **1b**.



Scheme 4 1,2-Shift mechanism of oxime–nitrone isomerization.



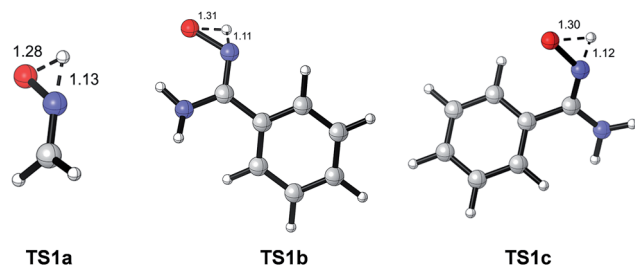
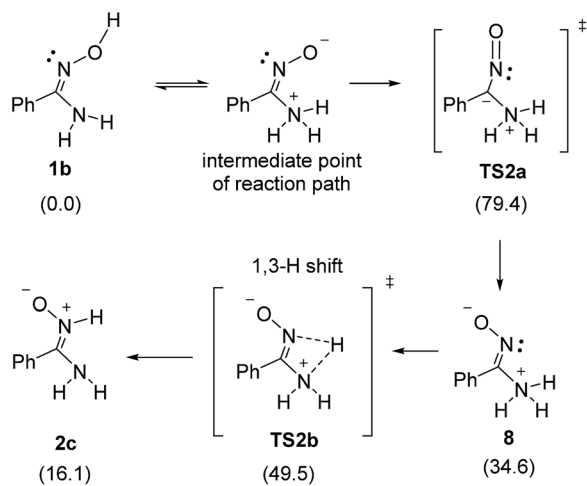


Fig. 2 Optimized transition structures (M06-2X/cc-pVTZ/PCM = EtOH) corresponding to 1,2-H shift between oximes **1** and nitrones **2**. Distances are given in angstrom.



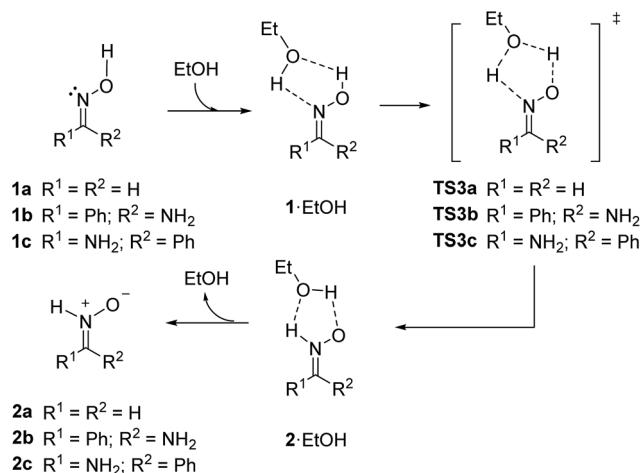
Scheme 5 1,2-H shift promoted by neighboring groups. Relative energies (M06-2X/cc-pVTZ/PCM = EtOH) are given in kcal mol⁻¹, between brackets.

shift (**TS2b**), to yield nitrone **2c** (for a complete analysis of the reaction including detailed IRCs and geometries of stationary points see ESI†). The first barrier (79.4 kcal mol⁻¹) corresponding to **TS2a** is too high to consider this mechanism as a possibility.

Next, we considered the possibility of bimolecular mechanisms. In the presence of a protic solvent as ethanol, a solvent molecule can transfer a proton to the nitrogen atom while receiving a different one from the oxime moiety, according to the mechanism illustrated in Scheme 6.

Similar results were obtained for the model system (series **a**) and for the real ones (series **b** and **c**). After formation of an initial complex between the oxime and a molecule of ethanol, promoted by H-bonding as depicted in Scheme 6, exchange of protons through **TS3** give rise to a new complex between ethanol and nitrone leading to isomerized nitrone **2**. The corresponding transition structures and the initial and final complexes formed with ethanol were calculated. The results are summarized in Table 3.

In the bimolecular process the formation of an initial complex should be stabilized by H-bonding. At the same time, the loss of translational entropy is much higher than in a unimolecular process; consequently, the value of free energy is



Scheme 6 Ethanol-catalyzed bimolecular isomerization of oxime **1** into nitrone **2** (M06-2X/cc-pVTZ/PCM = EtOH).

markedly overestimated.³³ Because of it, in order to compare the stability of reagents and products with initial and final complexes, it is more advisable to consider electronic energy (E_0) rather than free energy (G) where the entropic factor is introduced.³⁴ According to E_0 values, complexes **1**·EtOH are *ca.* 5–7 kcal mol⁻¹ more stable than the corresponding isolated oximes (5.5 kcal mol⁻¹ for **1a**, 6.5 kcal mol⁻¹ for **1b** and 7.0 kcal mol⁻¹ for **1c**). Complexes **2**·EtOH are about 7–9 kcal mol⁻¹ more stable than the parent isolated nitrones (6.8 kcal mol⁻¹ for **2a**, 9.2 kcal mol⁻¹ for **2b** and 9.3 kcal mol⁻¹ for **2c**).

Once formed the complex the isomerization takes place in the single entity oxime–EtOH to form the nitrone–EtOH entity, as in a unimolecular reaction, so the relative activation entropy

Table 3 Calculated (M06-2X/cc-pVTZ/PCM = EtOH) absolute (hartrees) and relative (kcal mol⁻¹) energies of the stationary points corresponding to isomerization of **1** into **2** promoted by a discrete molecule of EtOH

	ΔE_0^a	ΔG^a
1a	0.0	0.0
1a ·EtOH	−5.5	4.1
TS3a	16.1	27.1
2a ·EtOH	3.0	12.5
2a	9.8	10.0
1b	0.0	0.0
1b ·EtOH	−6.5	3.4
TS3b	13.2	24.5
2b ·EtOH	−0.1	10.0
2b	9.1	9.2
1c	6.2	5.8
1c ·EtOH	−0.8	9.1
TS3c	19.0	30.1
2c ·EtOH	6.6	16.8
2c	15.9	16.1

^a Referred to **1a** or **1b**; a discrete molecule of EtOH ($\Delta E_0 = -154.950425$ hartrees; $\Delta G = -154.975812$ hartrees) has been considered when necessary.

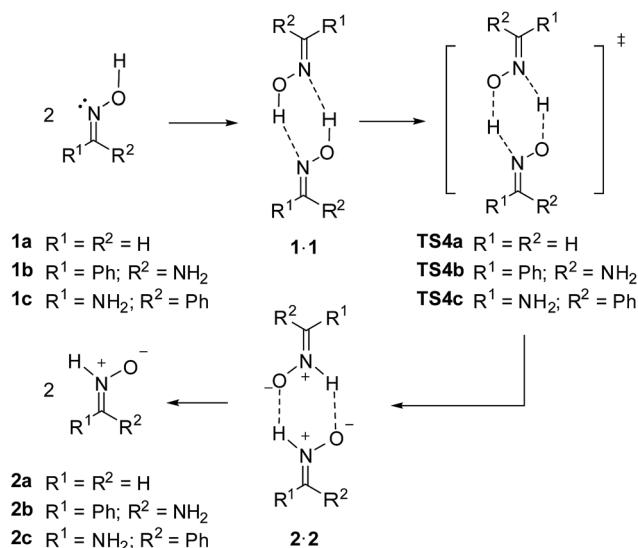


should be close to zero and Gibbs energy can be adequately considered for evaluating the barrier of the process. Once formed the initial complex **TS3** controls the rate of the reaction. Similar energy barriers were found in all cases (27.1 kcal mol⁻¹ for the model **1a/2a** and 24.5 kcal mol⁻¹ for the most stable *Z*-isomer (**1b/2b**) of the real system). The obtained values were 30 kcal mol⁻¹ lower than the unimolecular process discussed above. The geometry of the transition structures **TS3a–c** showed some differences between the model and the real system (Fig. 3). For **TS3a** similar distances were found between heteroatoms and hydrogens (1.26–1.28 Å) with the only exception of the shorter O_{EtOH}–H distance of 1.17 Å. In the case of **TS3b** and **TS3c** the distance O1–H (1.09 Å in both) was slightly shorter than the N2–H distance (1.12 Å in both). However, the distance between hydrogens and the oxygen atom of the ethanol molecule was much longer (1.41–1.47 Å). The spatial disposition of hydrogen atoms and heteroatoms were rather similar in all transition structures **TS3**. The higher stability of **TS3b** over **TS3c** is due to the extra H-bond interaction between O1 and the amino group.

The isomerization also takes place in aprotic solvents and under a variety of conditions. Consequently, it is plausible to consider, independently of the environment, a bimolecular mechanism in which only two molecules of oxime/nitrone are exclusively involved (Scheme 7). Again, formation of initial and final complexes should be considered. The corresponding transition structures and the initial and final complexes were calculated. The results are summarized in Table 4. The geometries of the transition structures are given in Fig. 4 (for geometries of complexes see ESI†).

By considering *E*₀ values as mentioned above, complexes **1·1** are more stable than the corresponding isolated oximes (8.6 kcal mol⁻¹ for **1a**, 11.1 kcal mol⁻¹ for **1b** and 5.3 kcal mol⁻¹ for **1c**). Complexes **2a·2a** and **2c·2c** are about 2 kcal mol⁻¹ more stable than the parent isolated nitrones (1.6 and 1.8 kcal mol⁻¹ for **2a** and **2c**, respectively). A greater difference (6.3 kcal mol⁻¹) was observed between complex **2b·2b** and the isolated nitrone **2b**. Again, the responsible for this difference is the H-bond interaction present in **2b**.

The isomerization from **1a·1a** to **2a·2a** is endergonic by 16.7 kcal mol⁻¹ with an energy barrier (**TS4a**) of 18.7 kcal mol⁻¹ from the ground state. In the case of the real system, the formation of two (*Z*)-nitrones from two molecules of (*Z*)-oxime is endergonic



Scheme 7 Bimolecular mechanism involving exclusively two molecules of oxime/nitrone.

by 14.5 kcal mol⁻¹. Calculations predicted a barrier (**TS4b**) of 15.3 kcal mol⁻¹ which is a lower value than those obtained for 1,2-H shift (79.4 kcal mol⁻¹), and intramolecular- and EtOH-assisted isomerizations (24.5 kcal mol⁻¹).³⁵ Similarly, if we consider the less stable (*E*)-isomers the isomerization from oximes to nitrones is endergonic by 16.8 kcal mol⁻¹ with an energy barrier (**TS4c**) of 16.4 kcal mol⁻¹.

The transition structure for the model isomerization of **1a** into **2a** is symmetrical with the hydrogen breaking and forming

Table 4 Calculated (M06-2X/cc-pVTZ/PCM = EtOH) absolute (hartrees) and relative (kcal mol⁻¹) energies of the stationary points corresponding to isomerization of **1** into **2** through a bimolecular mechanism

	ΔE_0^a	ΔG^a
1a	0.0	0.0
1a·1a	−8.6	1.4
TS4a	7.6	18.7
2a·2a	8.2	18.1
2a	9.8	10.0
1b	0.0	0.0
1b·1b	−11.1	0.0
TS4b	2.9	15.3
2b·2b	2.8	14.5
2b	9.1	9.2
1c	6.2	5.8
1c·1c	0.9	11.9
TS4c	20.1 ^b	28.2
2c·2c	17.7	28.7
2c	15.9	16.1

^a Referred to **1a** or **1b**; two molecules have been considered when necessary. ^b Barrier calculated from *E* values, without ZPVE correction. After ZPVE correction the value of the barrier is 16.7 kcal mol⁻¹ revealing the introduction of an experimental error that situates **2c·2c** above **TS4**. This error only affected **TS4c** since the rest of energy barrier values are similar after ZPVE correction.

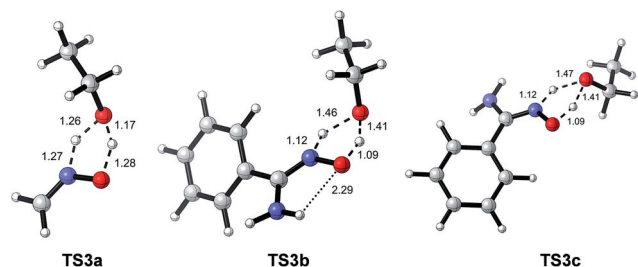


Fig. 3 Optimized transition structures (M06-2X/cc-pVTZ/PCM = EtOH) corresponding to isomerization of **1** into **2** promoted by a discrete molecule of EtOH. Distances are given in angstrom.



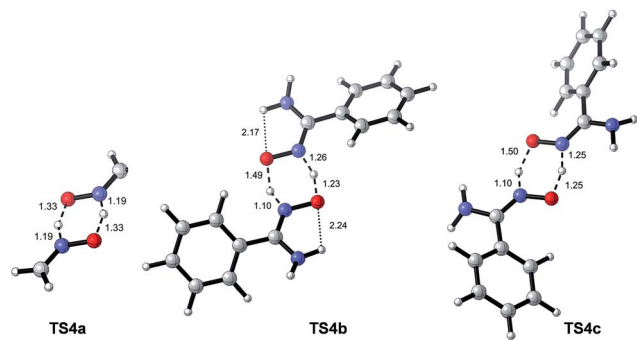


Fig. 4 Optimized transition structures (M06-2X/cc-pVTZ/PCM = EtOH) corresponding to isomerization of **1** into **2** through a bimolecular mechanism. Distances are given in angstrom.

bonds at distances of 1.19 Å and 1.33 Å from nitrogen and oxygen atoms, respectively. The transition state is completely planar without any deviation out of plane. On the other hand, for the real system the presence of the substituents distort the symmetrical disposition of the transition structure and an asynchronous H-exchange is observed. In fact, for **1b** several transition structures displaying minor conformational differences were located. The most stable **TS4b** and **TS4c** are given in Fig. 4. The shortest distance (1.10 Å in both **TS4b** and **TS4c**) corresponded to one forming N2–H bond, the other one having a distance of 1.26 Å for **TS4b** and 1.25 Å for **TS4c**. The longest distances correspond to the breaking O1–H bonds (1.49 Å and 1.50 Å in **TS4b** and **TS4c**, respectively) opposed to the above mentioned shortest N2–H forming bonds. The remaining O1–H breaking bonds showed distances of 1.23 Å and 1.25 Å in **TS4b** and **TS4c**, respectively. The hydrogen atoms and the heteroatoms involved in the proton exchange are all in the same plane without deviation. The only deviation out of the plane observed corresponded to the phenyl rings, already discussed above for both oximes and nitrones. Again, the difference in stability between **TS4b** and **TS4c** is attributed to a cooperative effect of the extra H-bonding interactions in **TS4b**. Such interactions are stronger (indicated by a shorter distance of 2.17 Å) in the side in which the hydrogen transfer towards the formation of the nitrone is more advanced.

In summary, these data with the simplest oxime **1a** and a real system **1b** corroborate that the oxime–nitron isomerization is a bimolecular process involving a hydrogen transfer from an oxime molecule to another. The lowest barrier (15.3 kcal mol^{−1}) for the isomerization has been found for the interconversion of two (*Z*)-oxime molecules into two (*Z*)-nitron molecules; starting from (*E*)-isomers the barrier was found to be 16.4 kcal mol^{−1}. Higher barriers were found for the other mechanisms investigated including unimolecular 1,2-shift (49.5 kcal mol^{−1}), neighboring group-assisted (79.4 kcal mol^{−1}) mechanisms, and bimolecular ethanol-assisted (24.5 kcal mol^{−1}) isomerization.

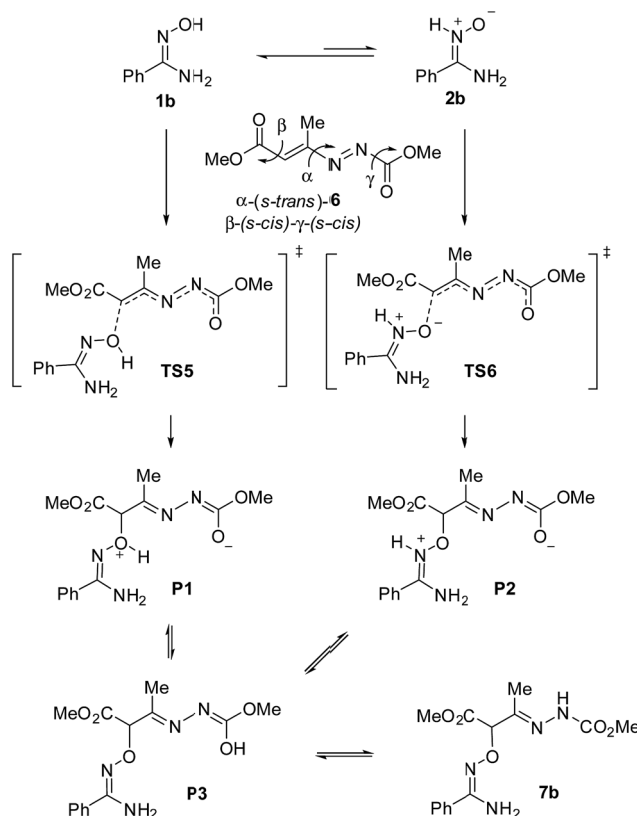
Michael addition of arylamidoximes to 1,2-diaza-1,3-dienes

As mentioned above, the reaction between an oxime and an electron-poor olefin, in the absence of a base, proceeds as an

ene-like reaction to yield nitrones (Scheme 2, path A). Interestingly, the reaction of oxime **1b** with 1,2-diaza-1,3-dienes such as **6**, in the absence of any base and in EtOH as a solvent, yields the corresponding *O*-alkylated oxime **7**,²¹ representing an exception to the usually observed behavior. 1,2-Diaza-1,3-dienes are well-known as Michael acceptors in β -position to the diazo functionality,³⁶ which is the directing electron-withdrawing group. This has been corroborated by reactivity indices calculations.³⁷ In order to investigate whether the nucleophilic addition to **6** takes place *via* the oxime **1b** or the nitron **2b** (after isomerization) we carried out a systematic computational study of the process in which we also included the corresponding (*E*)-isomers **1c** and **2c** (for a preliminary study with nitron **1a** see ESI†).

1,2-Diaza-1,3-dienes like **6** can adopt *s-trans* and *s-cis* conformations involving three dihedral angles. The most stable conformers (by *ca.* 4–5 kcal mol^{−1})³⁸ correspond to an *s-trans* conformation at the central dihedral angle.³⁹ However, the *s-cis* conformation might lead to stabilizing interactions that could furnish more stable situations. Because of it, both conformations have been taken into consideration. A comprehensive exploration of the PES of the reaction covering a total of 192 initial transition structures was carried out (for details see ESI†).

In principle, both the oxime **1b** and nitron **2b** can react with 1,2-diaza-1,3-diene **6** through the corresponding transition structures **TS5** and **TS6** to provide zwitterions **P1** and **P2**



Scheme 8 Mechanisms for the reaction between **1b** and **6**. Only (*Z*)-isomers are shown.



(Scheme 8). Further migration of the proton, through enol **P3**, yields the final product **7** observed experimentally. Such a migration might also occur during the formation of the new C–O bond, depending on the interactions found in the corresponding transition structures. Of course, it will be necessary to consider energetics of the isomerization discussed in the preceding section for the full analysis of the process. Also, *E/Z* isomerization of oxime and/or nitron tautomer could occur during the reaction. Because of it we have included such variability in the analysis as discussed above (not shown in Scheme 8).

The above-mentioned exploratory work of the PES for the nucleophilic attack provided, after full optimization at M06-2X/cc-pVDZ/PCM = EtOH level, 99 independent transition structures from which only 27 account for 100% of probability. Moreover, from those 27, only 8 contribute with more than 1%, accounting for 97.4% of probability. Notably, all TSs in the first 54 positions correspond to a nucleophilic attack of the nitron tautomer, the first 17 corresponding to (*Z*)-**2b**. The first instance of a transition structure corresponding to the nucleophilic attack of the oxime tautomer appears at 55th position with a relative free energy of 10.9 kcal mol^{−1} (for details see ESI†). Next, in order to obtain more accurate energy barriers for the reaction, we considered transition structures up to 4 kcal mol^{−1} of relative difference (19 TSs accounting for 99.8%) and they were full optimized at the highest triple- ζ level M06-2X/cc-pVTZ/PCM = EtOH. The results are given in Table 5. As mentioned above all transition structures correspond to the attack of nitron (*Z*)-**2b** (Table 5, entries 1–18) with one exception that

corresponds to (*E*)-**2c** (Table 5, entry 19), the most stable transition structure for the (*E*)-series. In order to compare the barriers of the different reaction pathways, we also included in triple- ζ optimization the most stable cases for the addition of oximes (*Z*)-**1b** and (*E*)-**1c** (Table 5, entries 20 and 21, respectively).⁴⁰

The three most stable transition states **TS6a–c** (within a range of 0.1 kcal mol^{−1}) and **TS6j** accounting for 64.7% correspond to the same approach; they only differ on variations at β and γ dihedral angles of the 1,2-diaza-1,3-diene **6**. The same situation is found for **TS6d**, **f**, **h** and **TS6e**, **g**, **TS6i** representing a different approach. In consequence, it is possible to limit the discussion by considering only 4 models, namely **TS6a**, **TS6d**, **TS6e** and **TS6i** that adequately represent 10 situations in a range of 2.0 kcal mol^{−1} of relative energy, accounting for 97.9% of Boltzmann's distribution. The rest of TSs, *i.e.* **TS6k–s**, are at least 2.0 kcal mol^{−1} above the most stable **TS6a** and account for the remaining 2.1%. The geometries of **TS6a**, **TS6d**, **TS6e** and **TS6i** are given in Fig. 5. Whereas in **TS6a** and **TS6i** the attack is carried out from the *Re* face of the nitron, in **TS6d** and **TS6e**, the attack comes from the *Si* face of the nitron. The forming bond distances are similar in all cases. **TS6a** and **TS6e** correspond to *droite* approaches with N2–O1–C1'–C2' angles of 44.1° and 13.0°, respectively. While **T6d** corresponds to a *gauche* approach, **TS6i** corresponds to an *anti* approach.

The four models illustrated in Fig. 5 are in a range of 1.8 kcal mol^{−1}, close to experimental error so, any of them could be considered valid. In any case, *the favored addition through the (Z)-nitron tautomer is fully confirmed*. The addition of (*E*)-

Table 5 Calculated (M06-2X/cc-pVTZ/PCM = EtOH) absolute (hartrees) and relative (kcal mol^{−1}) energies for the most stable transition structures of the reaction of oximes **1b**, **c** and nitrones **2b**, **c** with 1,2-diaza-1,3-diene **6**^a

Entry	Taut. ^b	TS	Face ^c	Stag. ^d	α^e	β^e	γ^e	ΔE_0^f	ΔG^f	Rel. ab. ^g (%)	Im. freq.
1	2b	TS6a	<i>Re</i>	g	t	c	c	0.0	0.0	23.4	−317.1
2		TS6b	<i>Re</i>	g	t	t	c	0.5	0.1	20.2	−304.1
3		TS6c	<i>Re</i>	g	t	c	t	0.4	0.1	20.2	−301.1
4		TS6d	<i>Si</i>	d	t	c	c	1.6	0.3	15.0	−331.7
5		TS6e	<i>Si</i>	g	t	c	c	1.4	0.6	8.6	−313.4
6		TS6f	<i>Si</i>	d	t	t	c	2.5	0.9	4.8	−321.6
7		TS6g	<i>Si</i>	g	t	c	t	2.4	1.5	1.9	−313.8
8		TS6h	<i>Si</i>	d	t	c	t	2.9	1.6	1.7	−330.8
9		TS6i	<i>Re</i>	a	t	c	c	3.3	1.8	1.1	−276.5
10		TS6j	<i>Re</i>	g	t	t	t	1.8	1.9	0.9	−297.2
11		TS6k	<i>Re</i>	g	c	c	c	4.0	2.0	0.8	−291.0
12		TS6l	<i>Si</i>	d	t	t	t	3.4	2.2	0.6	−317.3
13		TS6m	<i>Si</i>	g	t	t	t	3.7	2.6	0.3	−318.4
14		TS6n	<i>Re</i>	a	t	t	c	4.2	2.7	0.3	−277.4
15		TS6o	<i>Si</i>	d	c	c	c	4.3	3.4	0.1	−321.6
16		TS6p	<i>Re</i>	g	c	c	t	5.1	3.8	0.0	−296.0
17		TS6q	<i>Si</i>	d	c	t	c	4.9	4.0	0.0	−321.4
18		TS6r	<i>Si</i>	d	c	c	t	5.1	4.0	0.0	−323.2
19	2c	TS6s	<i>Re</i>	g	t	c	c	5.1	5.0	0.0	−254.6
20	1b	TS5a	<i>Re</i>	a	t	c	c	14.2	13.2	0.0	−386.4
21	1c	TS5b	<i>Re</i>	a	c	t	c	19.5	19.4	0.0	−552.8

^a Only those below 4 kcal mol^{−1} of relative free energy in the 2 ζ study have been considered for the triple- ζ one (see text). ^b Referred to the tautomer that reacts with **6**. ^c Indicates the face of the oxime/nitron. ^d Three staggered orientations of the oxime/nitron attack to **6**, defined as “d” (*droite*), “g” (*gauche*) and “a” (*anti*); see ESI for definition of d/g/a. ^e “c” and “t” correspond to *s-cis* and *s-trans* conformations of the considered dihedral angles of **6**. ^f With respect to the most stable one. ^g According to Boltzmann's distribution.



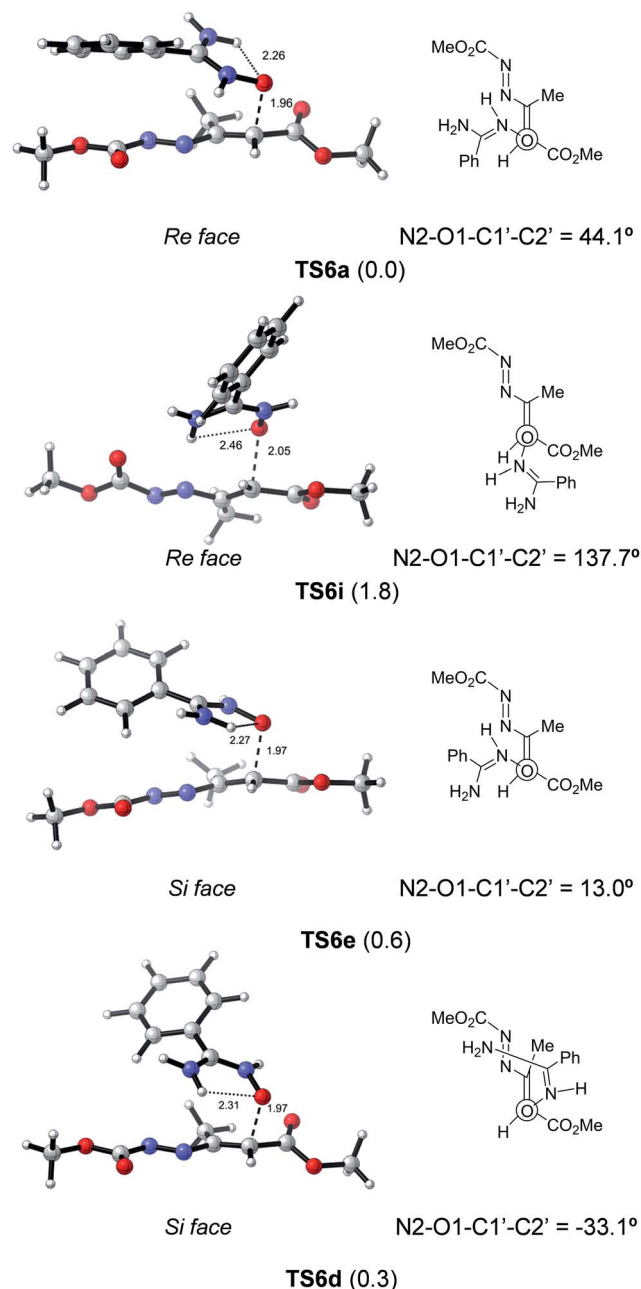


Fig. 5 Selected optimized transition structures (M06-2X/cc-pVTZ/PCM = EtOH) corresponding to the nucleophilic attack of **2b** to **6**. The more stable TSs for each approach are shown. Relative energies are given in kcal mol⁻¹. Distances are given in angstroms.

nitron **2c** was found to be 5.0 kcal mol⁻¹ higher in energy. The most stable TSs, **TS5a** and **TS5b**, corresponding to the addition of oxime tautomers gave even higher energy values (by 13.2 and 19.4 kcal mol⁻¹ for (*Z*)-oxime **1b** and (*E*)-oxime **1c**, respectively).

Fig. 6 illustrates the compared simplified energy diagrams for the different isomers showing the energy barriers for the reaction of each isomer. Different final points (not shown in Fig. 6) were obtained for each reaction path; after tautomeric processes those points led to either **7b** or **7c**, the most stable tautomers (for complete energy diagrams, transition structures and full reaction paths see ESI†).

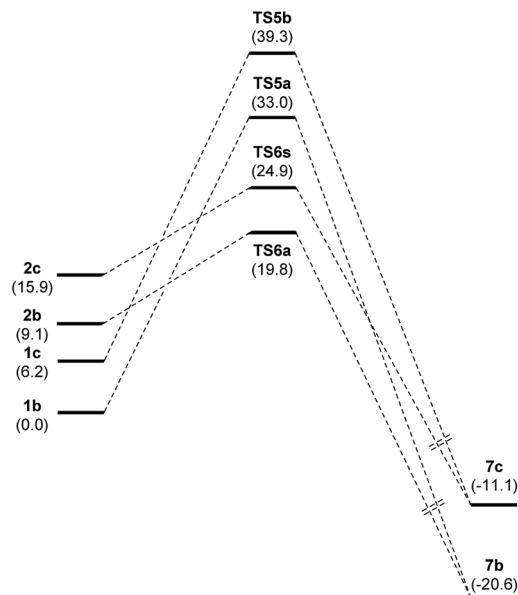


Fig. 6 Simplified free energy diagrams (M06-2X/cc-PVTZ/PCM = EtOH) for the nucleophilic addition of oximes **1b**, **c** and nitrones **2b**, **c** to 1,2-diaza-1,3-diene **6**. **TS5a** and **TS5b** correspond to (*Z*)- and (*E*)-oximes, respectively. **TS6a** and **TS6b** correspond to (*Z*)- and (*E*)-nitrones, respectively.

To sum up, even though isomerization of oxime **1b** to nitron **2b** has a barrier of 15.3 kcal mol⁻¹ and requires a cost in energy of 9.2 kcal mol⁻¹, the most stable transition state is that corresponding to the attack of the nitron tautomer through **TS6a**. According to the Curtin-Hammett principle⁴¹ this is the preferred path for the reaction, revealing a rare case in which a nitron tautomer participates in a nucleophilic addition reaction.

Ene-like reactions and alternative processes

As we discussed in the introduction, ene-like reaction between **1b** and **6** (path A, Scheme 2) and dipolar cycloaddition between **2b** and **6** (path D, Scheme 2), could be competitive with the nucleophilic Michael addition. In order to validate our model and further confirm the preference by the nitron-mode addition, we also calculated those alternative paths of the reaction.

The possibility of an ene-like reaction (Scheme 2, path A) was first explored. A typical transition structure **TS_A** could be located in a simple model like addition of **1b** to dimethyl 2-methyl-fumarate.⁴² The IRC analysis corroborated that the reaction was concerted and transfer of hydrogen takes place giving rise to the corresponding nitron (see ESI†).

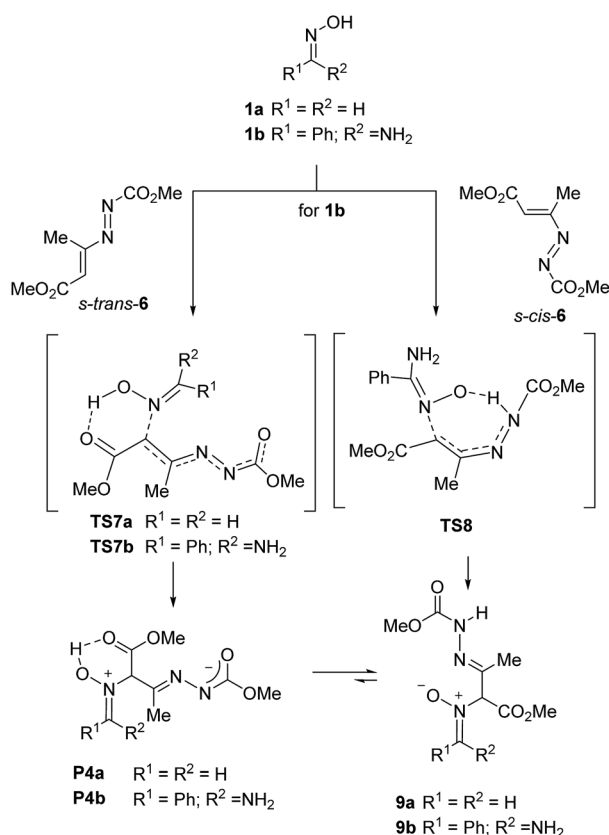
On the other hand, the reaction with 1,2-diaza-1,3-diene **6** led to dramatic changes on the located transition structures. A full exploration of the PES corresponding to the nucleophilic attack of the oxime nitrogen was carried out by considering **1b** and **1c** as well as *s-trans*-**6** and *s-cis*-**6** (for details see ESI†).

No transition structures corresponding to an H-transfer to the carbon atom through a concerted ene-like process could be located; instead, in all cases the hydrogen atom was oriented to the formation of an H-bond either with one of the nitrogen



atoms of the diazo group or the carbonyl ester group directly linked to the double bond. The most stable paths were those showed in Scheme 9. **TS7b** was 0.6 kcal mol⁻¹ lower in energy than **TS8** (energy barriers: 20.6 and 21.2 kcal mol⁻¹ for **TS7b** and **TS8**, respectively) so, both transition structures can be considered competitive. Transition structure **TS7b** does not correspond to an ene-like process and the IRC analysis confirmed a partial hydrogen transfer. The resulting zwitterion **P4b** is assumed to be transformed into the neutral nitron **9b** through a typical tautomeric process. In the case of *s-cis*-**6**, IRC of **TS8** reflects a complete transfer of hydrogen to the diazo group, leading directly to the neutral nitron. Any attempt of locating an ene-like transition structure similar to that located for the model methyl 2-methylfumarate, led to one of the TSs located previously during PES exploration (see ESI†).

The free energy barriers found for **TS7b**, and **TS8** were 0.8 and 1.4 kcal mol⁻¹, respectively, higher in energy than the most stable **TS6a** (Fig. 7). These values indicate that the nucleophilic addition is preferred in agreement with experimental observations. However, it should also be considered that both processes might be competitive since the differences observed between barrier values are within the error limit of the calculations. Nevertheless, it would not be surprising that a close inspection of the reaction mixture could reveal the obtention of minor amounts of nitron **9b**.



Scheme 9 Preferred path for the nucleophilic attack of the oxime nitrogen.

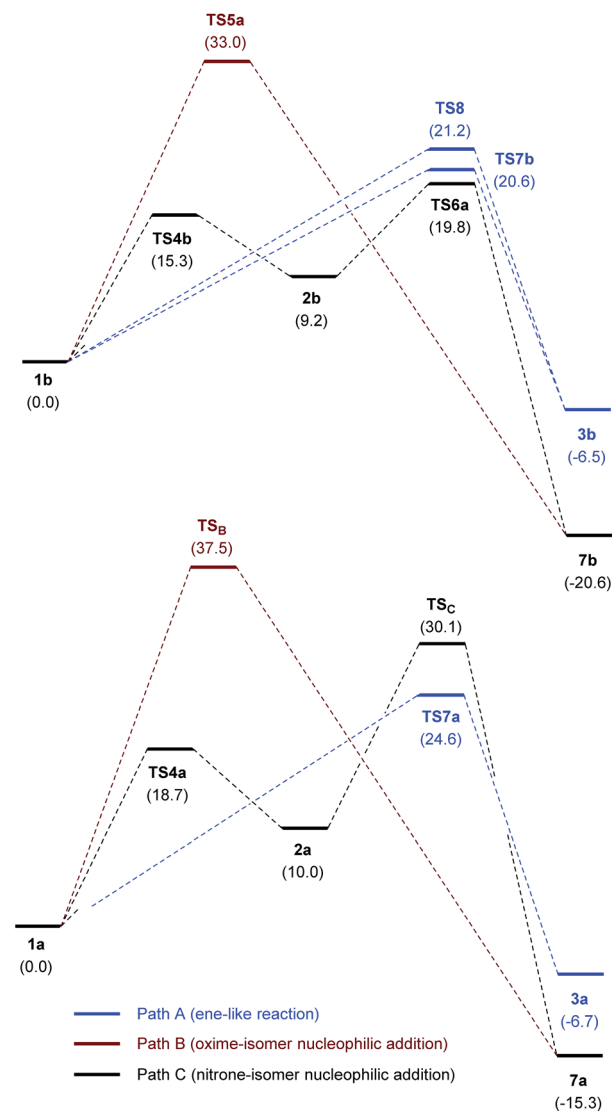


Fig. 7 Comparative free energy diagrams (M06-2X/cc-pVTZ/PCM = EtOH) for the reaction between **1b** and **6** (top) and **1a** and **6** (bottom). Relative free energies are given in kcal mol⁻¹.

When the same study was made with the unsubstituted oxime **1a**, the nucleophilic addition of the oxime nitrogen leading to the nitron through **TS7a**, clearly resulted the most stable situation (5.5 kcal mol⁻¹ lower in energy than the corresponding transition structure **TS8**; see ESI†) in agreement with previous experimental findings with aldoximes and revealing the particularity of oxime **1b** (Fig. 7). These results allow predicting that, in general and in the absence of a base, the ene-like process (Scheme 2, path A) should be preferred for the reaction of an oxime and an electron-poor alkene, but when steric hindrance is present as in the case of ketoximes like **1b**, the *O*-alkylation is a competitive process taking place through the attack of the nitron tautomer (Scheme 2, path C).

The reason of the different results obtained with **1a** and **1b** can be found in the different spatial requirements of both oximes. The geometries of transition structures **TS7b**, **TS8** and **TS7a** are given in Fig. 8. The unsubstituted oxime **1a** attacks



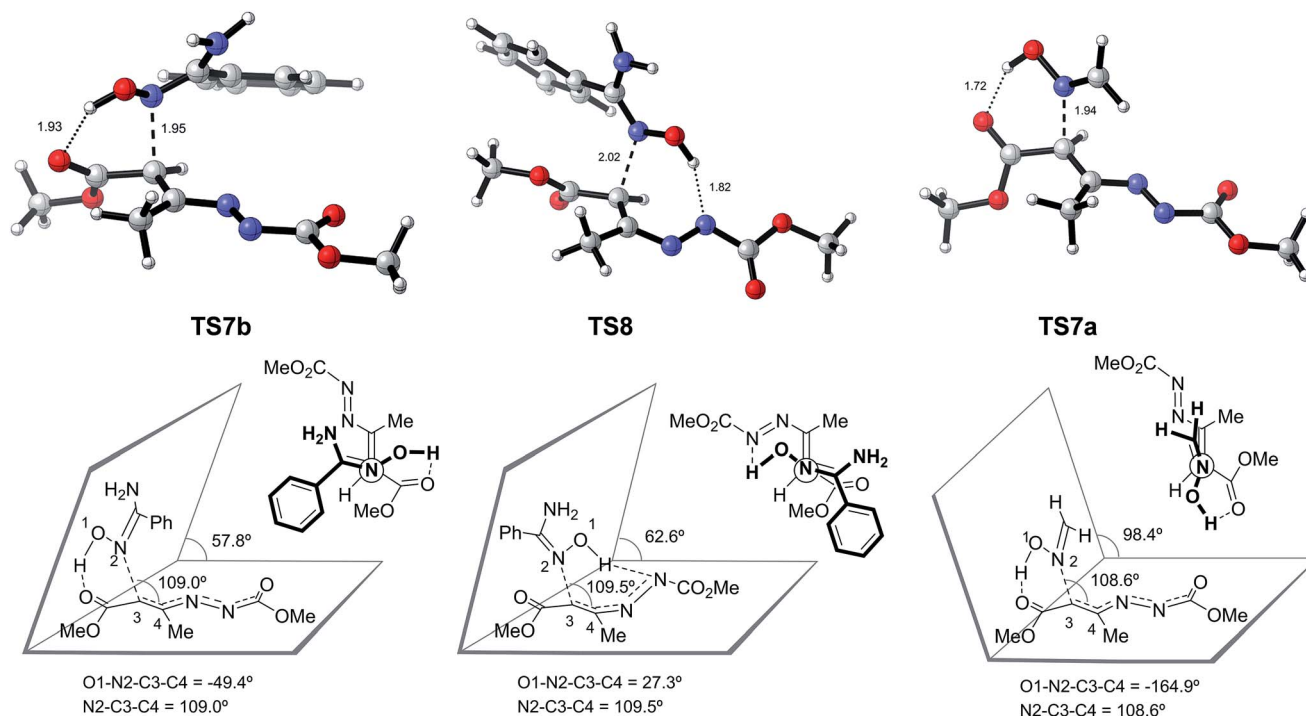


Fig. 8 Optimized transition structures (M06-2X/cc-pVTZ/PCM = EtOH) of **TS7b** and **TS8** corresponding to the ene-like reaction between oxime **1b** and 1,2-diaza-1,3-diene **6**, and **TS7a** corresponding to the reaction between **1a** and **6**. Distances are given in angstrom.

with a correct orientation of the nitrogen lone pair, through a completely perpendicular eclipsed orientation (O1-N2-C3-C4 dihedral angle of -164.9°) following a typical Burgi-Dunitz trajectory (N2-C3-C4 angle of 108.6°). The perpendicularity of the oxime with respect to the 1,2-diaza-1,3-diene is evidenced by the angle of 98.4° formed between the reactant planes defined by the oxime and the diazadiene (Fig. 8).⁴³ On the other hand, oxime **1b** is forced to modify such orientation in order to maintain the Burgi-Dunitz trajectory (N2-C3-C4 angles of 109.0° and 109.5° for **TS7b** and **TS8**, respectively) and, at the same time, to avoid unfavorable steric interactions. As a consequence, the nitrogen lone pair is not perfectly aligned with the carbon atom and the situation increases the energy of the corresponding transition structures. The distortion during the attack of **1b** is revealed by the steeper angles formed between the planes defined by the oxime and the diazadiene (57.8° and 62.6° for **TS7b** and **TS8**, respectively) with respect to that observed for **1a**.

Finally, no concerted transition structures corresponding to path D (Scheme 2) could be located. Moreover, any attempt of locating such concerted transition structures led to some of the already characterized TSs listed in Table 5. In particular, a systematic search and optimization for initial 3,5-*endo* and 3,5-*exo* transition structures led to **TS6d** and **TS6a**, respectively.

Conclusions

The oxime-nitrone isomerization has been studied computationally and it has been determined that it takes place through a bimolecular mechanism involving two molecules of oxime (or

nitrone). The commonly accepted 1,2-H shift can be completely discarded, even if favored by neighboring groups, due to the high energies of the corresponding transition structures. Oximes usually react with electron deficient olefins to give the corresponding nitrones through *N*-alkylation (formal Michael addition) and concomitant hydrogen transfer. The mechanism of the reaction had been proposed to be an ene-like concerted process and it has been confirmed in the case of a simple model.

The *O*-alkylation observed experimentally in the reaction of oxime **1b** to 1,2-diaza-1,3-diene **6**, in the absence of a base, can be considered an exception. The computational study carried out on this process has demonstrated that the reaction takes place through the nucleophilic attack of the nitrone tautomer. The addition represents the first example in which the nitrone tautomer is involved into a nucleophilic addition of an oxime. Nevertheless, the *N*-alkylation should be considered a competitive process. Indeed, it can also be predicted that *O*-alkylation (though nitrone tautomer) will take place particularly with sterically demanding substrates such as ketoximes like the studied **1b**. In the case of substrates that allow a favorable orthogonal approach of the oxime to the electron deficient alkene, the *N*-alkylation leading to the corresponding nitrone will be preferred.

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