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Cite this: *RSC Adv.*, 2014, 4, 32415

A new C–C bond formation model based on the quantum chemical topology of electron density†

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ELF topological analyses of bonding changes in non-polar, polar and ionic organic reactions involving the participation of C=C(X) double bonds make it possible to establish a unified model for C–C bond formation. This model is characterised by a C-to-C coupling of two *pseudoradical* centers generated at the most significant atoms of the reacting molecules. The global electron density transfer process that takes place along polar and ionic reactions favours the creation of these *pseudoradical* centers at the most nucleophilic/electrophilic centers of the reacting molecules, decreasing activation energies. The proposed reactivity model based on the topological analysis of the changes in electron density throughout a reaction makes it possible to reject the frontier molecular orbital reactivity model based on the analysis of molecular orbitals.

Received 8th May 2014
 Accepted 2nd July 2014

DOI: 10.1039/c4ra04280h

www.rsc.org/advances

Establishing organic molecular mechanisms based on quantum chemistry calculations and the transition state theory

From the advance of numerical computation at the end of the 20th century, computational quantum chemistry (CQC) has

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† Dedicated to Prof B. Silvi for his contribution to the development of the ELF topological analysis of electron density in the study of structure and reactivity.



Luis R. Domingo obtained his PhD in Organic Chemistry in 1987. In 1990 he became an Associate Professor, and in 2010 obtained his current position of Full Professor at the Department of Organic Chemistry of the University of Valencia. In 1995 he came up with his first studies in the field of Theoretical Organic Chemistry. So far, he has published more than 220 theoretical studies in different fields of

Organic Chemistry. His research interest comprises the study of molecular mechanisms of cycloaddition reactions, the study of organocatalysis, and the application of the DFT reactivity indices in Organic Chemistry. He says: while the distribution of the electron density is responsible for the molecular shape and physical properties, the capability for changes in electron density is responsible for the reactivity.

been greatly accepted by organic chemists due to its practicality in the study of reaction mechanisms involving actual molecules with 50–70 atoms. CQC enables the localisation and characterisation of the reagents, products, transition state structures (TS) and intermediates involved in organic reactions, thus making the study of molecular mechanisms of organic reactions possible. The comparison of computed activation parameters obtained from the transition state theory (TST)^{1–3} with those experimentally obtained by kinetic experiments allows performing an analysis of available computational models. Feasible competitive reaction channels can be theoretically studied, and thus, explain experimental outcomes.

Two appealing data are first obtained through CQC in a straightforward manner: (i) total electronic energies; and (ii) molecular geometries. Although electronic energies associated with the stationary points involved in a chemical reaction are very dependent on the computational level, geometries are less dependent on it. Thus, while *ab initio* Hartee–Fock (HF) calculations⁴ widely used along the two last decades of the 20th century allowed the obtention of good TS geometries, that theoretical level yielded very high activation energies overestimating TS energies. Consequently, very time-consuming post-HF energy calculations⁴ were performed related to HF optimised geometries. The development of the density functional theory (DFT)⁵ at the end of the 20th century, whose calculations provided activation energies closer to experimental values, allowed the standardisation of DFT computations in the study of organic reactions. Thus, several DFT functionals such as B3LYP,^{6,7} MPWB1K⁸ and more recently M06-2X,⁹ which provide accurate energies, have been developed, allowing the study of organic reaction with a computational demand similar to HF calculations.



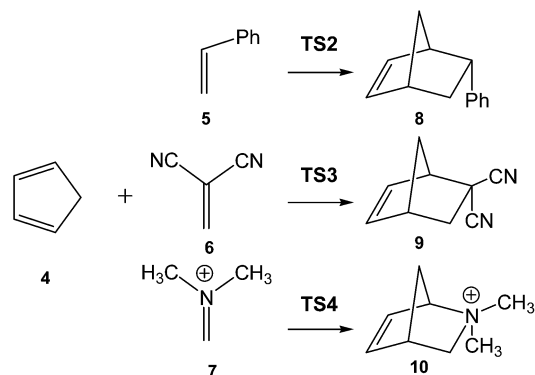


Scheme 2 DA reactions of Cp 4 with ethylenes of increased electrophilic character.

bond at the asynchronous TSs were found in the narrow range from 2.17 to 1.96 Å. In order to analyse some significant characteristics of the C–C bond formation in DA reactions, three different reactions, *i.e.* the DA reactions of Cp 4 with styrene 5, with 1,1-dicyanoethylene (DCE) 6, and with iminium cation 7, have been selected (see Scheme 3). Relative energies and TS geometries are given in Fig. 2.

Interestingly, as can be seen in Fig. 2, while the relative energies associated with these DA reactions were found in a wide range from 21 to -5 kcal mol^{-1} , the TS geometries were found to be very similar; the three TSs showed an asynchronous C–C single bond formation. If the most unfavourable TS2 is ruled out, TS3 and TS4 are found to be highly asynchronous. On the other hand, the three TSs showed a similar C–C single bond formation, the distances between the two interacting carbons at the TSs being in the narrow range from 2.04 to 1.96 Å. The C–C distances were only 0.2 Å shorter than the distances found at the very unfavourable synchronous TS1. Unfortunately, no information about the evolution of the bonding changes in the reactions can be obtained from these energy- and geometrical parameters; thus, with this information the nature of the molecular mechanism remains unresolved.

The results obtained from numerous theoretical studies devoted to DA reactions, pose an important question: what is the origin for the different activation energies found in TSs having similar geometries? The response was obtained after



Scheme 3 DA reactions of Cp 4 with styrene 5, with DCE 6, and with iminium cation 7.



Fig. 2 TSs involved in the DA reactions of Cp 4 with styrene 5, TS2, with dicyanoethylene 6, TS3, and with iminium cation 7, TS4. Relative energies are given in kcal mol^{-1} , while distances are given in Angstroms.

analysing a large number of theoretical studies devoted to cycloaddition reactions.

In 1999, in an earlier paper, the DA reactions of nitroethylene 11 with three ethylenes of increased electron-rich character were studied (see Scheme 4).¹⁴ A good correlation between the activation energies of the reactions and the nucleophilic/electrophilic behaviours of the reagents was established. The increase of the electron-rich character of the ethylene, *i.e.* its nucleophilic character, goes together with a decrease of the activation energy associated with the C–C bond formation.

This finding was quantitatively ascertained to analyse the electrophilicity ω index¹⁵ of the reagents involved in DA reactions.¹⁶ This first DFT study enabled the establishment of a unique scale of electrophilicity, in which both dienes and dienophiles were included,¹⁶ thus permitting the establishment of a good correlation between the difference of the electrophilicity ω indices of the reagents, $\Delta\omega$, and the feasibility of the DA reactions.¹⁶ The more electrophilic a reagent, *i.e.* when it is located at the top of the scale, and more nucleophilic the other reagent, *i.e.* when it is located at the bottom of the table, the more polar and faster the reaction.

Later, in 2003 the DA reactions between Cp 4 and the cyanoethylene series 14, experimentally studied by Sauer in 1964 (ref. 17) and given in most organic textbooks as an example of the effects of the electron-withdrawing substitution in DA reactions,¹⁸ were analysed (see Scheme 5).¹⁹

Some appealing conclusions were obtained from that DFT study: (i) both synchronous and asynchronous TSs were found in this series of DA reactions, depending on the symmetric

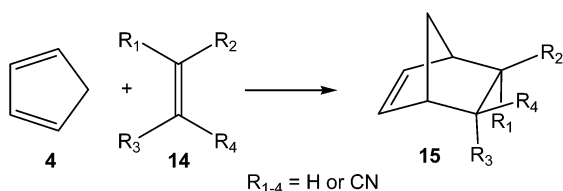


Scheme 4 DA reactions of nitroethylene 11 with three ethylenes of increased nucleophilic character.



substitution of the ethylene; *i.e.* while the symmetrically substituted 1,2-dicyanoethylenes maleonitrile (**14a**, $R_1 = R_3 = \text{CN}$, $R_2 = R_4 = \text{H}$) and fumaronitrile (**14b**, $R_1 = R_4 = \text{CN}$, $R_2 = R_3 = \text{H}$) yielded synchronous TSs, the asymmetric substituted DCE **6** (**14c**, $R_1 = R_2 = \text{CN}$, $R_3 = R_4 = \text{H}$) yielded a highly asynchronous TS;¹⁹ (ii) the synchronicity in the C–C single bond formation appears to be an unfavourable factor, *i.e.* the DA reaction with DCE **6** is *ca.* 1000 times faster than that with symmetric maleonitrile **14a** and fumaronitrile **14b**; and (iii) interestingly, a very good correlation between the global electron density transfer (GEDT) found at the TSs and the logarithm of the experimental rate constant was established, indicating that the GEDT could be one of the key factors in the activation energy (see Fig. 3).¹³ The GEDT at the TSs is computed by sharing the natural charges at the TSs obtained by natural bond orbital (NBO) analysis^{20,21} between the nucleophilic and the electrophilic frameworks.¹³ The GEDT concept comes from the observation that the electron density transfer that takes place along polar and ionic reactions is not a local process, but a global flux of electron density taking place from the nucleophile to the electrophile, and not being dependent on the approach mode of both reagents.²²

These results made it possible to establish the polar Diels–Alder (P-DA) mechanism,¹³ in which favourable electrophilic/nucleophilic interactions along the polar reaction are responsible for the GEDT found at the TSs, and consequently, for the feasibility of the reaction. The proposed polar mechanism for



Scheme 5 DA reactions between Cp **4** and cyanoethylene series **14**.



Fig. 3 Plot of the logarithm of the experimental rate constant k vs. GEDT, in e , $R^2 = 0.99$, for the DA reactions of Cp **4** with ethylene **2**, and the cyanoethylene series.

DA reactions has two very significant repercussions: (i) most DA reactions do not take place through a pericyclic mechanism²³ as proposed in all textbooks,^{24–26} (ii) DA reactions are not a special type of organic reactions. Note that most of the DA reactions taking place through highly asynchronous TSs present a C–C bond formation similar to that found in most polar organic reactions.

From these findings, the different activation energies of the DA reactions given in Scheme 3 can be explained. The computed GEDT values at the TSs given in Fig. 2 are $0.06e$ at **TS2**, $0.28e$ at **TS3** and $0.45e$ at **TS4**. Although only three reactions are dealt with herein, a very good correlation between the GEDT and the computed activation energy can be established ($R^2 = 0.98$, see Fig. 4). Note that when the aforementioned twelve DA reactions are considered, this correlation remains at $R^2 = 0.89$.¹³ The lineal correlation between polarity of the reaction, measured by the GEDT at the TS, and the corresponding activation energy, allowed the establishment of an appealing classification of DA reactions into non-polar Diels–Alder (N-DA) reactions, characterised by a GEDT below $0.2e$, and P-DA reactions characterised by a GEDT above $0.2e$.¹³ While N-DA reactions are of little synthetic interest as they demand harsh reaction conditions, the feasibility of a P-DA reaction increases with the polar character of the reaction; *i.e.* the electrophilic/nucleophilic character of the reagents. Interestingly, these behaviours can easily be anticipated by analysing the electrophilicity ω (ref. 15) and the nucleophilicity N indices,^{27,28} defined within the conceptual DFT,^{29,30} at the ground state of the reagents. A good correlation between the activation energies of the DA reactions of Cp **4** with ethylenes of different electron-withdrawing substitution shown in Scheme 2 and the global electrophilicity ω indices was found, $R^2 = 0.92$.¹³ Similarly, a good correlation between the logarithm of the reaction rate constant of 5-substituted indoles with a series of benzhydryl cations, experimentally studied by Mayr,³¹ with the global nucleophilicity N indices has also been established, $R^2 = 0.98$.²⁸



Fig. 4 Plot of the activation energies ΔE , in kcal mol^{-1} , vs. GEDT, in e , for the DA reactions given in Scheme 3.



Of the twelve DA reactions studied, the fastest one was that between Cp **4**, a nucleophilic neutral molecule, and the iminium cation **7**, a very high electrophilic cationic species (see Scheme 3). To distinguish the P-DA reactions in which the polarity is only evidenced in the course of the reaction from those reactions in which all species are ionic, the latter reaction was classified as an ionic Diels–Alder (I-DA) reaction.¹³

What is the origin of the GEDT in polar and ionic organic reactions? An appealing concept defined within DFT is the electronic chemical potential μ (ref. 32) which correlates with the absolute electronegativity χ through the simple equation $\chi = -\mu$.³³ The electronic chemical potential μ , defined as $\mu = (\partial E/\partial N)_v$, is associated with the feasibility to exchange electron density of a molecule with the environment at ground state.³² Sanderson suggested the electronegativity equalisation principle³⁴ in chemistry, in which electronegativity tends to equalise. The correctness of Sanderson's principle immediately comes from the fact that the electronic chemical potential μ is a property of an equilibrium state. Consequently, when two molecules A and B, with $\mu_A < \mu_B$, approach one another, there is a flux of electron density from A, the less electronegative species, towards B, the more electronegative one, to equilibrate the electronic chemical potential μ_{AB} in the new interacting system. The larger the electronic chemical potential difference, $\Delta\mu_{A-B}$, the larger the GEDT. It is noteworthy that in P-DA reactions taking place along high asynchronous TSS, the maximum GEDT value is reached after passing the TS, and when the first C–C single bond is formed. In experimental stepwise P-DA reactions,^{35,36} this value is reached with the formation of the corresponding zwitterionic intermediate.

In spite of this appealing finding above-mentioned, some issues remained unresolved: how do the bonding changes in both N-DA and P-DA reactions take place? and, how can the GEDT modify the pattern in bonding changes along the reaction? That is, how can the polarity of the reaction modify the reaction mechanism? To answer these questions a complete analysis of the bonding changes along a reaction must be performed. This task demands a complete quantum topology analysis of the electron density obtained from the quantum wave function along the IRC of an elemental reaction.

Establishing organic reaction mechanisms based on the quantum chemical topology analysis of electron density

Since the introduction of the chemical bond concept by G. N. Lewis at the beginning of the 20th century³⁷ many theoretical models have been developed to understand matter structure and chemical reactivity. Quantum chemical tools based on the valence bond (VB) theory,^{38–40} the molecular orbital (MO) theory,⁴ and most recently DFT⁵ have proven to be useful to chemists. However, in spite of the advances made in this field, the characterisation of chemical bonds, and more specifically

the breaking/forming processes along a reaction, appear to be unresolved.⁴¹ Like many other chemical concepts, chemical bonds are defined in a rather ambiguous manner as they are not observable, but rather belong to a representation of the matter at a microscopic level which is not fully consistent with quantum mechanical principles.

To harmonise the chemical description of matter with quantum chemical postulates, several mathematical models have been developed. Among them, the theory of dynamical systems,⁴² convincingly introduced by Bader through the theory of atoms in molecules (AIM),⁴³ has become a powerful method of analysis. The AIM theory enables a partition of the electron density within the molecular space into basins associated with atoms. The development of the AIM theory was the origin of a significant contribution to conceptual chemistry in the definition of concepts such as the atom inside a molecule or bond critical points.^{44–46} For many years, Popelier has been studying the AIM quantum chemical topology (QCT) of the electron density for characterising chemical bonds, having performed a great deal of work in different fields such as the characterisation of heterocyclic rings,⁴⁷ atomic properties of aminoacids,⁴⁸ pK_a predictions,⁴⁹ and radicals.⁵⁰

Another appealing procedure that provides a more straightforward connection between the electron density distribution and the chemical structure is the QCT analysis of the electron localisation function (ELF) of Becke and Edgecombe.⁵¹ In this sense, Silvi and Savin presented the ELF in a very chemical fashion, using their topological analysis as an appealing model of chemical bonding.^{52–55} After an analysis of the electron density, ELF divides the electron density of a molecule into basins, *i.e.* domains in which the probability of finding an electron pair is maximal. Basins are classified as core basins and valence basins. The latter are characterised by the synaptic order, *i.e.* the number of atomic valence shells in which they participate.⁵⁶ Thus, there are monosynaptic, disynaptic, trisynaptic basins and so on. Monosynaptic basins, labelled $V(A)$, correspond to lone pairs or non-bonding regions, while disynaptic basins, labelled $V(A, B)$, connect the core of two nuclei A and B and, thus, correspond to a bonding region between A and B. This description recovers the Lewis bonding model, providing a very suggestive graphical representation of the molecular system. Analysis of the ELF valence basin populations N provides similar GEDT values of than those obtained by NBO analysis.^{22,57}

On the other hand, the characterisation of the electron density reorganisation to evidence the bonding changes along a reaction path is the most attractive method to characterise a reaction mechanism.^{58–60} To quantitatively perform these analyses, the bonding evolution theory (BET), consisting of the joint-use of ELF topology and Thom's catastrophe theory^{61–63} (CT) was proposed by Krokidis *et al.*⁶⁴ as a new tool for analysing the electronic changes in chemical processes, being applied to different elementary reactions.^{65–78} In this field, Andrés performed a systematic investigation characterising the mechanisms of significant organic reactions such as DA reactions,^{58,79} [3 + 2] cycloaddition (32CA) reactions,⁸⁰ the Bergman cyclisation,⁸¹ the Cope rearrangement,⁸² and the Nazarov cyclisation.⁸³



ELF topological model for the C–C bond formation in non-polar reactions

In 2003, the molecular mechanism of the DA reaction between butadiene **1** and ethylene **2**, given in Scheme 1, was characterised using BET.⁵⁸ Two appealing conclusions were obtained from this QCT analysis of the bonding changes along the one-step mechanism: (i) the reaction consists of seven differentiated phases characterised by 10 catastrophes, within of the classification given by Thom,⁸⁴ belonging to the fold and cusp types. Each one of these phases is characterised by a bonding change in the Lewis structure with respect to the previous phase; (ii) the formation of the C–C single bond in this DA reaction takes place through a C-to-C coupling between two *pseudoradical* centers^{85,86} generated through the breaking of the C=C double bonds, at a C–C distance of 2.04 Å (see Fig. 5).

The seven phases characterising this N-DA reaction have recently been categorised into four groups associated with significant chemical bonding changes (see Fig. 6).⁸⁷ (i) in the first one, A (in red colour), the three C=C double bonds present in butadiene **1** and ethylene **2** break, phases I to III; (ii) in the second group, B (in green colour), the formation of two *pseudoradical* structures takes place by gathering electron density at the end carbons of the two unsaturated reagents, phases IV and V (see the four V(C) monosynaptic basins, integrating *ca.* 0.5e each one, in structure **16** in Fig. 5). The electron density demanded for the creation of the *pseudoradical* centers comes from the depopulation of the C=C double bonds of the butadiene and ethylene moieties; (iii) in the third group, C (in blue colour), which is constituted only by the most relevant phase VI, the formation of the two new C–C single bonds takes place through the C-to-C coupling between the *pseudoradical* centers generated in the previous group B (see the two disynaptic basins V(Cx, Cy), integrating *ca.* 1.0e each one, in structure **17** in Fig. 5) (iv) in the fourth group, D (in violet colour), while the formation of the two C–C single bonds is completed, the formation of the new C=C double bonds takes place at the end of the IRC. This QCT analysis of the bonding changes along the one-step mechanism of the N-DA reaction between butadiene **1** and ethylene **2** shows that the bonding changes are non-concerted.⁸⁷



Fig. 5 Most relevant ELF attractors in the structures of phase V, structure **16**, and phase VI, structure **17**, involved in the synchronous C–C single bond formation in the N-DA reaction between butadiene **1** and ethylene **2**. The *pseudoradical* centers involved in the C–C bond formation are characterised by the four V(C) monosynaptic basins present in structure **16**.

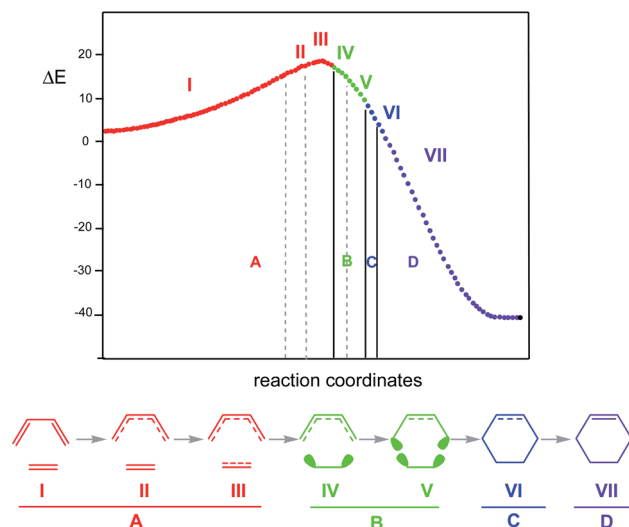


Fig. 6 Reaction path calculated by means of the IRC method for the DA reaction between butadiene **1** and ethylene **2**. ΔE are in kcal mol⁻¹. A bonding between atoms in all phases is demonstrated by the standard Lewis representation; however, in the case of phases IV and V, ellipses reflect the non-bonding electron density concentrated in the C atoms. The four colours represent the four main groups in which the bonding changes can be categorised.

Later, an ELF topological comparative study between the one-step and stepwise mechanisms of the N-DA of Cp **4** with ethylene **2** was performed in order to characterise these competitive molecular mechanisms (see Scheme 6).⁸⁸ As expected, BET analysis of the one-step mechanism of the N-DA reaction of Cp **4** with ethylene **2** was found to be very similar to that found in the N-DA reaction of butadiene **1** with ethylene **2**.⁵⁸ Again, ELF topological analysis showed that the synchronous C–C single bond formation takes place at a C–C distance of 1.84 Å by the C-to-C coupling of two *pseudoradical* centers generated in Cp **4** and in ethylene **2**. Interestingly, a similar pattern for the C–C single bond formation along the first step of the stepwise mechanism was found.⁸⁸ In the stepwise mechanism, the formation of the first C–C single bond takes place also by a C-to-C coupling between two radical centers, yielding the formation of



Scheme 6 One-step and stepwise pathways of the N-DA reaction between Cp **4** and ethylene **2**.



the diradical intermediate **18** (see Scheme 6). In this very unfavourable step, the C–C bond formation begins at a distance of 1.77 Å.

Some appealing conclusion was obtained from this comparative ELF topological analysis: (i) although the one-step mechanism and the stepwise mechanism appear to have a dissimilar electron reorganisation, note that the one-step mechanism is still classified as a pericyclic reaction,²³ topologically, they present many similarities. The C–C bond formation along the one-step mechanism takes place by coupling of two *pseudoradical* centers (see *pseudodiradical* structure **20** in Fig. 7), while along the stepwise mechanism it takes place by coupling of two radical centers (see the diradical structure **21** in Fig. 7).⁸⁸ Note that while the *pseudodiradical* structures are obtained by B3LYP restricted calculations to have a closed shell, the *diradical* structures such as **18** and **21** must be obtained by UB3LYP unrestricted calculations to have an open shell; (ii) ELF topological analysis of bonding changes along N-DA reactions of butadiene **1** and Cp **4** with ethylene **2**, showed the non-concerted nature of breaking and forming bonds along these symmetric cycloaddition reactions, making it possible to reject the pericyclic mechanism for these reactions.⁸⁷

An analysis of bonding changes taking place along the energy profiles of N-DA reactions, allowed explaining the high activation energy required in non-polar processes. In all the studied reactions involving closed-shell molecules, the formation of the C–C single bonds takes place after passing the TS. The proposed model for the C–C bond formation in non-polar reactions suggests that it takes place by merging the electron density of two carbons having non-bonding electron density, which comes from the depopulation of electron density of the bonding region of the C=C double bonds present in these reagents. ELF topological analysis of the corresponding *pseudodiradical* structures indicates that at this stage of the reaction the C=C double bonds are already broken. Consequently, the high activation energy associated with non-polar reactions can be related to the unfavourable energy associated with the breaking of the C=C double bond in these non-concerted processes.⁸⁸



Fig. 7 Semblance between the *pseudodiradical* structure **20** involved in the one-step mechanism, and the diradical structure **21** involved in the stepwise mechanism of the N-DA reaction between Cp **4** and ethylene **2**.

Several studies devoted to the characterisation of the molecular mechanism of non-polar reactions have supported this model for the C–C bond formation in non-polar processes. Thus, an ELF analysis of the electron reorganisation along the non-polar 32CA reaction of carbonyl ylide **22** with tetramethylethylene **23** (see Scheme 7) showed that the C–C bond formation in this 32CA reaction takes place at a C–C distance of 2.18 Å by a C-to-C coupling of two *pseudoradical* centers.⁸⁹

Interestingly, carbonyl ylide **22** already has a *pseudodiradical* character at its ground state (see structure **22** in Scheme 7). This behaviour accounts for the very low activation energy found in this 32CA reaction,^{89,90} and for the advanced character of the C–C bond formation. Note that organic reactions involving open-shell radical species are fast.

Recently, an ELF analysis for the C–C bond formation step in the N-heterocyclic carbene (NHC) catalysed hydroacylation of unactivated C–C double bonds (see Scheme 8) showed that in this non-polar reaction the C–C bond formation takes place also by a C-to-C coupling of a *pseudodiradical* structure **27** generated at the C1 and C2 carbons along the non-polar reaction (see Fig. 8).⁹¹

On the other hand, BET analysis along the cyclisation reactions of 1,3-butadiene **1**,⁹² 1,3,5-hexatriene **28**,⁹² and 1,3,5,7-octatetraene **29** (ref. 93) (see Scheme 9) showed that the C–C bond formation in these electrocyclic reactions also takes place by a C-to-C coupling of two *pseudoradical* centers generated at the end of these conjugated systems.

Very recently, Andrés studied the cycloheptatriene **33**/norcaradiene **34** isomerisation (see Scheme 10).⁶⁰ As in the isomerisation of hexatriene **28**, this reaction is an electrocyclic reaction allowing the formation of a new C–C single bond between the end carbons of the hexatriene system of **33**. The authors commented “once the TS has been reached, two new monosynaptic basins appear at the end carbons of the hexatriene system. This phase is very short, immediately these two



Scheme 7 32CA reaction of carbonyl ylide **22** with tetramethylethylene **23**.



Scheme 8 The C–C bond formation step in the NHC catalysed hydroacylation of unactivated C–C double bonds.





Fig. 8 Pseudoradical structure 27 generated along the C–C bond formation step in the N-heterocyclic carbene-catalysed hydroacylation of unactivated C–C double bonds.



Scheme 9 Electrocyclic reactions of the conjugated polyenic systems 1, 28 and 29.

monosynaptic basins merge into the disynaptic $V(C1, C6)$ basin accounting for the closing of the cyclopropane ring⁶⁰ (see the merger of the $V(C1)$ and $V(C6)$ monosynaptic basins present in structure 35 into the $V(C1, C6)$ disynaptic basin present in structure 36 in Scheme 10).

a) cycloheptatriene 33 / norcaradiene 34 isomerisation.



b) C–C bond formation by coupling of the two pseudoradical centers.



Scheme 10 Cycloheptatriene 33/norcaradiene 34 isomerisation reaction.

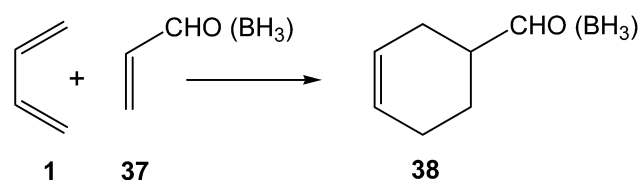
ELF topological model for the C–C bond formation in polar reactions

After performing a BET analysis of the N-DA of butadiene 1 with ethylene 2, the molecular mechanism of the P-DA reactions between butadiene 1 and acrolein 37, in the absence and the presence of a Lewis acid catalyst, BH_3 , were investigated in 2006 by using BET (see Scheme 11).⁷⁹ These P-DA reactions take place by a one-step mechanism. BET analysis of the bonding changes along the two P-DA reactions indicated that they take place along eleven, in the non-catalysed reaction, and ten, in the LA-catalysed reaction, differentiated phases. In P-DA reactions, taking place along highly asynchronous TSSs, the formation of the two C–C single bonds takes place in two differentiated phases of the IRC, thus characterising these reactions by a *two-stage one-step* mechanism.⁹⁴ Interestingly, the formation of the first C–C single bond in these P-DA reactions showed a similar pattern to that found in the N-DA reaction between butadiene 1 and ethylene 2; *i.e.* the formation of the first C–C single bond takes place after passing the TSs, at a C–C distance of 1.92 Å, by a C-to-C coupling of two *pseudoradical* centers, in both non-catalysed and LA-catalysed reactions.⁷⁹

Later, an ELF topological study on the origin of the synchronicity in bond formation in the P-DA reactions of Cp 4 with DCE 6, and with tetracyanoethylene (TCE) (14d, $R_1 = R_3 = R_2 = R_4 = CN$) established that, as the N-DA reaction between Cp 4 and styrene 5 or ethylene 1,⁸⁸ the C–C single bond formation in these polar reactions takes place also by a C-to-C coupling of two *pseudoradical* centers generated along the reaction (see Fig. 9).⁹⁵

Interestingly, as in the P-DA reaction between butadiene 1 and acrolein 37, ELF analysis of the C–C bond formation at the P-DA reaction of Cp 4 with DCE 6 showed that the formation of the first C–C single bond in the *two-stage one-step* mechanism takes place at a C–C distance of 1.95 Å through the most favourable two-center interaction between the one of the most nucleophilic centers of Cp 4 and the most electrophilic center of DCE 6.⁹⁵ This favourable interaction, which is responsible for the high regioselectivity experimentally observed in most P-DA reactions,^{22,96} is in agreement with the creation of the two *pseudoradical* centers at the most nucleophilic center of Cp 4 and the most electrophilic center of DCE 6 (see structure 39 in Fig. 9).

ELF topological studies of DA reactions have shown that while in N-DA reactions the electron density demanded for the formation of the *pseudoradical* centres is reached mainly by the



Scheme 11 P-DA reactions between butadiene 1 and acrolein 37, in the absence and the presence of a Lewis acid catalyst.





Fig. 9 ELF attractors at selected points of the IRC of the P-DA reactions between Cp 4 and DCE 6, 39 and 40, and TCE 14d, 41 and 42.

depopulation of the C=C double bonds present in the reagents,⁸⁸ in P-DA reactions this electron density mainly comes from the GEDT that takes place along the polar process. Thus, the *pseudoradical* center created in the electrophilic species is formed mainly in the most electrophilic center of the molecule, which is the center with the highest spin density achieved through the GEDT process.⁹⁷ These findings allowed proposing, within the conceptual DFT, first the nucleophilic P_k^- and electrophilic P_k^+ Parr functions,^{98,99} and later the radical P_k^0 Parr functions¹⁰⁰ in order to characterise the most relevant reactive centers within an organic molecule (see Fig. 10). Parr functions provide comparable information to that obtained using Fukui functions,¹⁰¹ but they are conceptually different; Fukui functions are approximated through the HOMO and LUMO frontier molecular orbitals (FMOs),¹⁰² while Parr functions are based on the changes of atomic spin density associated to GEDT processes.⁹⁸

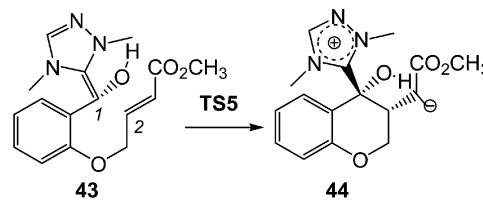
This model for the C-C bond formation in polar reactions has been supported by a number of ELF topological studies



Fig. 10 Nucleophilic P_k^- function of styrene 5, electrophilic P_k^+ Parr function of DCE 6, and radical P_k^0 Parr functions of iminium cation 7.

devoted to the characterisation of the molecular mechanism in diverse polar reactions. Thus, an ELF topological analysis for the C-C bond formation step in the intramolecular Stetter reaction¹⁰³ (see Scheme 12) established that it takes place also by a C-to-C coupling of two *pseudoradical* centers generated at the most nucleophilic center of Breslow intermediate 43, the C1 carbon, and the most electrophilic center of the α,β -unsaturated ester framework of 43, the C2 carbon. Interestingly, TS5 already showed the presence of the *pseudoradical* centers at the C1 and the C2 carbons (see Fig. 11). Note that TS5, which was associated with a Michael-type addition, presents a similar topology to that found in the non-polar structure 27.

A very recent ELF topological analysis for the C-C bond formation step in the NHC catalysed Michael addition of enols to α,β -unsaturated acyl-azoliums (see Scheme 13) showed that it takes place also at a C-C distance of 2.00 Å by a C-to-C coupling of two *pseudoradical* centers generated at the most nucleophilic center of enol 45, the C1 carbon, and the most electrophilic center of the α,β -unsaturated acyl-azoliums 46, the conjugated C2 carbon.¹⁰⁴



Scheme 12 The C-C bond formation step in the intramolecular Stetter reaction.



Fig. 11 TS5 associated with the C-C bond formation in Breslow intermediate 43.



Scheme 13 The C-C bond formation step in the NHC catalysed Michael addition of enols to α,β -unsaturated acyl-azoliums.



- 25 F. A. Carey and R. J. Sundberg, *Advanced Organic Chemistry. Part A: Structure and Mechanisms*, Springer, New York, 2007.
- 26 I. Fleming, *Molecular Orbitals and Organic Chemical Reactions*, Wiley, 2009.
- 27 L. R. Domingo, E. Chamorro and P. Pérez, *J. Org. Chem.*, 2008, **73**, 4615.
- 28 L. R. Domingo and P. Pérez, *Org. Biomol. Chem.*, 2011, **9**, 7168.
- 29 P. Geerlings, F. De Proft and W. Langenaeker, *Chem. Rev.*, 2003, **103**, 1793.
- 30 D. H. Ess, G. O. Jones and K. N. Houk, *Adv. Synth. Catal.*, 2006, **348**, 2337.
- 31 S. Lakhdar, M. Westermaier, F. Terrier, R. Goumont, T. Boubaker, A. R. Ofial and H. Mayr, *J. Org. Chem.*, 2006, **71**, 9088.
- 32 R. G. Parr and W. Yang, *Density Functional Theory of Atoms and Molecules*, Oxford University Press, New York, 1989.
- 33 R. G. Parr, R. A. Donnelly, M. Levy and W. E. Palke, *J. Chem. Phys.*, 1978, **68**, 3801.
- 34 R. T. Sanderson, *Science*, 1955, **121**, 207.
- 35 V. Y. Korotaev, A. Y. Barkov, P. A. Slepukhin, M. I. Kodess and V. Y. Sosnovskikh, *Mendeleev Commun.*, 2011, **21**, 112.
- 36 S. Lakhdar, F. Terrier, D. Vichard, G. Berionni, N. El Guesmi, R. Goumont and T. Boubaker, *Chem.-Eur. J.*, 2010, **16**, 5681.
- 37 G. N. Lewis, *J. Am. Chem. Soc.*, 1916, **38**, 762.
- 38 J. C. Slater, *Phys. Rev. B: Solid State*, 1931, **37**, 481.
- 39 J. C. Slater, *Phys. Rev. B: Solid State*, 1931, **38**, 1109.
- 40 L. Pauling, *Nature*, 1948, **161**, 1019.
- 41 B. T. Sutcliffe, *Int. J. Quantum Chem.*, 1996, **58**, 645.
- 42 R. H. Abraham and C. D. Shaw, *Dynamics: The Geometry of Behavior*, Addison-Wesley, Redwood City, CA, 1992.
- 43 R. F. W. Bader, *Atoms in Molecules. A Quantum Theory*, Clarendon Press, Oxford, UK, 1990.
- 44 R. F. W. Bader, S. G. Anderson and A. J. Duke, *J. Am. Chem. Soc.*, 1979, **101**, 1389.
- 45 R. F. W. Bader, T. T. Nguyendang and Y. Tal, *J. Chem. Phys.*, 1979, **70**, 4316.
- 46 R. F. W. Bader, *Theor. Chem. Acc.*, 2001, **105**, 276.
- 47 M. Griffiths and P. Popelier, *J. Chem. Inf. Model.*, 2013, **53**, 1714.
- 48 P. Popelier and F. Aicken, *ChemPhysChem*, 2003, **4**, 824.
- 49 A. Harding, D. Wedge and P. Popelier, *J. Chem. Inf. Model.*, 2009, **49**, 1914.
- 50 N. Singh, P. Popelier and P. O'Malley, *Chem. Phys. Lett.*, 2006, **426**, 219.
- 51 A. D. Becke and K. E. Edgecombe, *J. Chem. Phys.*, 1990, **92**, 5397.
- 52 A. Savin, A. D. Becke, J. Flad, R. Nesper, H. Preuss and H. G. von Schnering, *Angew. Chem., Int. Ed.*, 1991, **30**, 409.
- 53 B. Silvi and A. Savin, *Nature*, 1994, **371**, 683.
- 54 A. Savin, B. Silvi and F. Colonna, *Can. J. Chem.*, 1996, **74**, 1088.
- 55 A. Savin, R. Nesper, S. Wengert and T. F. Fassler, *Angew. Chem., Int. Ed.*, 1997, **36**, 1808.
- 56 B. Silvi, *J. Mol. Struct.*, 2002, **614**, 3.
- 57 L. R. Domingo, P. Pérez and J. A. Sáez, *RSC Adv.*, 2013, **3**, 7520.
- 58 S. Berski, J. Andrés, B. Silvi and L. R. Domingo, *J. Phys. Chem. A*, 2003, **107**, 6014.
- 59 V. Polo, J. Andrés, S. Berski, L. R. Domingo and B. Silvi, *J. Phys. Chem. A*, 2008, **112**, 7128.
- 60 J. Andrés, P. González-Navarrete and V. Safont, *Int. J. Quantum Chem.*, 2014, DOI: 10.1002/qua.24665.
- 61 R. Thom, *Structural Stability and Morphogenesis: An Outline of a General Theory of Models*, Inc.:Reading, MA, 1976.
- 62 A. E. R. Woodcock and T. Poston, *A Geometrical Study of Elementary Catastrophes*, Spinger-Verlag, Berlin, 1974.
- 63 R. Gilmore, *Catastrophe Theory for Scientists and Engineers*, Dover, New York, 1981.
- 64 X. Krokidis, S. Noury and B. Silvi, *J. Phys. Chem. A*, 1997, **101**, 7277.
- 65 S. Berski and Z. Latajka, *Int. J. Quantum Chem.*, 2011, **111**, 2378.
- 66 S. Berski, F. R. Sensato, V. Polo, J. Andrés and V. S. Safont, *J. Phys. Chem. A*, 2011, **115**, 514.
- 67 E. Chamorro, J. C. Santos, B. Gomez, R. Contreras and P. Fuentealba, *J. Chem. Phys.*, 2001, **114**, 23.
- 68 E. Chamorro, J. C. Santos, B. Gomez, R. Contreras and P. Fuentealba, *J. Phys. Chem. A*, 2002, **106**, 11533.
- 69 P. Chaquin and A. Scemama, *Chem. Phys. Lett.*, 2004, **394**, 244.
- 70 D. B. Chesnut and L. J. Bartolotti, *Chem. Phys.*, 2000, **257**, 175.
- 71 I. Fourre, B. Silvi, P. Chaquin and A. Sevin, *J. Comput. Chem.*, 1999, **20**, 897.
- 72 F. Fuster, A. Sevin and B. Silvi, *J. Phys. Chem. A*, 2000, **104**, 852.
- 73 N. Gillet, R. Chaudret, J. Contreras-Garcia, W. T. Yang, B. Silvi and J. P. Piquemal, *J. Chem. Theory Comput.*, 2012, **8**, 3993.
- 74 X. Krokidis, V. Goncalves, A. Savin and B. Silvi, *J. Phys. Chem. A*, 1998, **102**, 5065.
- 75 X. Krokidis, N. W. Moriarty, W. A. Lester and M. Frenklach, *Chem. Phys. Lett.*, 1999, **314**, 534.
- 76 I. M. Ndassa, B. Silvi and F. Volatron, *J. Phys. Chem. A*, 2010, **114**, 12900.
- 77 V. Polo, P. González-Navarrete, B. Silvi and J. Andrés, *Theor. Chem. Acc.*, 2008, **120**, 341.
- 78 J. P. Salinas-Olvera, R. M. Gómez and F. Cortés-Gúzman, *J. Phys. Chem. A*, 2008, **112**, 2906.
- 79 S. Berski, J. Andrés, B. Silvi and L. R. Domingo, *J. Phys. Chem. A*, 2006, **110**, 13939.
- 80 V. Polo, J. Andrés, R. Castillo, S. Berski and B. Silvi, *Chem.-Eur. J.*, 2004, **10**, 5165.
- 81 J. C. Santos, J. Andrés, A. Aizman, P. Fuentealba and V. Polo, *J. Phys. Chem. A*, 2005, **109**, 3687.
- 82 V. Polo and J. Andrés, *J. Comput. Chem.*, 2005, **26**, 1427.
- 83 V. Polo and J. Andrés, *J. Chem. Theory Comput.*, 2007, **3**, 816.
- 84 R. Thom, *Stabilité Structurale et Morphogénèse*, Paris, 1972.
- 85 L. A. Errede, J. M. Hoyt and R. S. Gregorian, *J. Am. Chem. Soc.*, 1960, **53**, 5224.



- 86 L. R. Domingo, E. Chamorro and P. Pérez, *Lett. Org. Chem.*, 2010, **432**.
- 87 L. R. Domingo, *Org. Chem. Curr. Res.*, 2013, **2**, 120.
- 88 L. R. Domingo, E. Chamorro and P. Pérez, *Org. Biomol. Chem.*, 2010, **8**, 5495.
- 89 L. R. Domingo and J. A. Sáez, *J. Org. Chem.*, 2011, **76**, 373.
- 90 L. R. Domingo and S. R. Emamian, *Tetrahedron*, 2014, **70**, 1267.
- 91 L. R. Domingo, J. A. Saéz and M. Arnó, *RSC Adv.*, 2012, **2**, 7127.
- 92 J. Andrés, S. Berski, L. R. Domingo, V. Polo and B. Silvi, *Curr. Org. Chem.*, 2011, **15**, 3566.
- 93 J. Andrés, S. Berski, L. R. Domingo and P. González-Navarrete, *J. Comput. Chem.*, 2012, **33**, 748.
- 94 L. R. Domingo, J. A. Saéz, R. J. Zaragoza and M. Arnó, *J. Org. Chem.*, 2008, **73**, 8791.
- 95 L. R. Domingo, P. Pérez and J. A. Sáez, *Org. Biomol. Chem.*, 2012, **10**, 3841.
- 96 L. R. Domingo, P. Pérez, M. J. Aurell and J. A. Sáez, *Curr. Org. Chem.*, 2012, **16**, 2343.
- 97 L. R. Domingo, M. J. Aurell, P. Pérez and J. A. Sáez, *RSC Adv.*, 2012, **2**, 1334.
- 98 L. R. Domingo, P. Pérez and J. A. Sáez, *RSC Adv.*, 2013, **3**, 1486.
- 99 E. Chamorro, P. Pérez and L. R. Domingo, *Chem. Phys. Lett.*, 2013, **582**, 141.
- 100 L. R. Domingo and P. Pérez, *Org. Biomol. Chem.*, 2013, **11**, 4350.
- 101 R. G. Parr and W. Yang, *J. Am. Chem. Soc.*, 1984, **106**, 4049.
- 102 P. W. Ayers, C. Morell, F. De Proft and P. Geerlings, *Chem.–Eur. J.*, 2007, **13**, 8240.
- 103 L. R. Domingo, R. J. Zaragoza, J. A. Saéz and M. Arnó, *Molecules*, 2012, **17**, 1335.
- 104 L. R. Domingo, J. A. Sáez and M. Arnó, *Org. Biomol. Chem.*, 2014, **12**, 895.
- 105 L. R. Domingo, M. J. Aurell and P. Pérez, *RSC Adv.*, 2014, **4**, 16567.
- 106 L. R. Domingo and P. Pérez, *Phys. Chem. Chem. Phys.*, 2014, **16**, 14108.
- 107 L. R. Domingo, *J. Org. Chem.*, 2001, **66**, 3211.
- 108 K. Fukui, *Molecular Orbitals in Chemistry Physics and Biology*, New York, 1964.

